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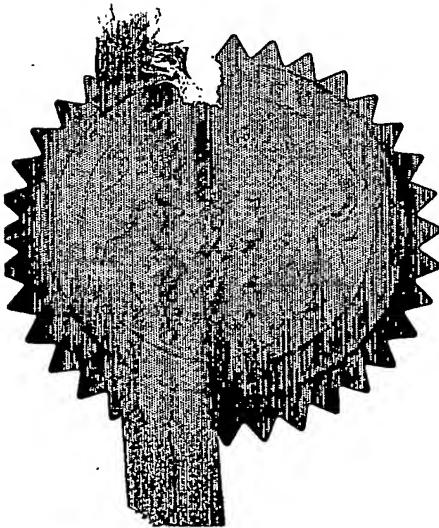
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1. Your reference	4-32572P1	0215844.2		
2. Patent application number (The Patent Office will fill in this part)	09 JUL 2002	10 JUL 02 E732004 1 D00524 P01/7700 0.00-0215844.2		
3. Full name, address and postcode of the or of each applicant (underline all surnames)	NOVARTIS AG LICHTSTRASSE 35 4056 BASEL SWITZERLAND	SWITZERLAND		
071925487005 Patent ADP number (if you know it) If the applicant is a corporate body, give the country/state of its incorporation				
4. Title of invention	Organic compounds			
5. Name of your agent (if you have one) "Address for service" in the United Kingdom to which all correspondence should be sent (including the postcode)	B.A. YORKE & CO. CHARTERED PATENT AGENTS COOMB HOUSE, 7 ST. JOHN'S ROAD ISLEWORTH MIDDLESEX TW7 6NH			
Patents ADP number (if you know it)	1800001	✓		
6. If you are declaring priority from one or more earlier patent applications, give the country and the date of filing of the or of each of these earlier applications and (if you know it) the or each application number	Country	Priority application number (if you know it)	Date of filing (day/month/year)	
7. If this application is divided or otherwise derived from an earlier UK application, give the number and the filing date of the earlier application	Number of earlier application	Date of filing (day/month/year)		
8. Is a statement of inventorship and of right to grant of a patent required in support of this request? (Answer 'Yes' if: a) any applicant named in part 3 is not an inventor, or b) there is an inventor who is not named as an applicant, or c) any named applicant is a corporate body. (see note (d))	Yes			

Patents Form 1/77

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Continuation sheets of this form

Description	49
Claim(s)	8
Abstract	
Drawing(s)	3m

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Priority documents

Translations of priority documents

Statement of inventorship and right to grant of a patent (Patents Form 7/77)

Request for preliminary examination and search (Patents Form 9/77)

Request for substantive examination (Patents Form 10/77)

Any other documents
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11.

I/We request the grant of a patent on the basis of this application

Signature

Date

B. A. Yorke & Co.

09 July 2002

12. Name and daytime telephone number of person to contact in the United Kingdom

Mrs. E. Cheetham
020 8560 5847

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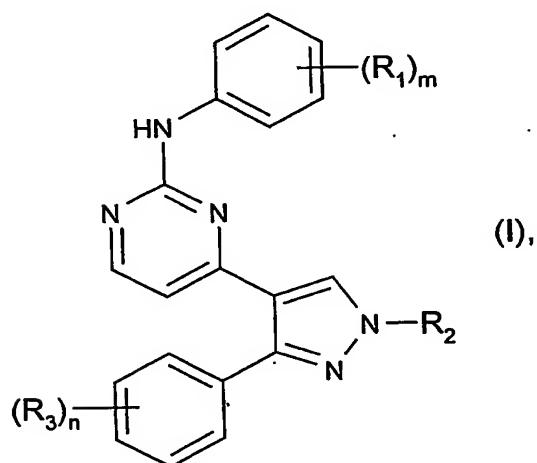
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Organic Compounds

The invention relates to phenyl-[4-(3-phenyl-1H-pyrazol-4-yl)-pyrimidin-2-yl]-amine derivatives and to processes for the preparation thereof, to pharmaceutical compositions comprising such derivatives and to the use of such derivatives – alone or in combination with one or more other pharmaceutically active compounds – for the preparation of pharmaceutical compositions for the treatment especially of a proliferative disease, such as a tumour.

The invention relates to phenyl-[4-(3-phenyl-1H-pyrazol-4-yl)-pyrimidin-2-yl]-amine derivatives of formula I



wherein

m and n are independently of one another an integer from 1 to 5;

R₁ is unsubstituted, mono- or di-substituted amino-sulfonyl; unsubstituted, mono- or di-substituted amino; a heterocyclic radical; lower alkyl substituted by a heterocyclic radical; a radical R₄-lower alkyl-X-, wherein R₄ is hydrogen, unsubstituted, mono- or di-substituted amino, or a heterocyclic radical, and X is -S- or -O-; or a radical R₅-C(=O)-, wherein R₅ is hydrogen, unsubstituted or substituted lower alkyl, free or etherified hydroxy, unsubstituted, mono- or di-substituted amino, or a heterocyclic radical; wherein the R₁ substituents are selected independently of one another if m>1;

or two vicinal R₁ substituents together with the phenyl carbon atoms to which they are attached form a heterocyclic ring;

R₂ is hydrogen, unsubstituted or substituted lower alkyl or a heterocyclic radical; and

R_3 is halogen; unsubstituted or substituted lower alkyl; or free, etherified or esterified hydroxy; wherein the R_3 substituents are selected independently of one another if $n > 1$; or two vicinal R_3 substituents together with the phenyl carbon atoms to which they are attached form a heterocyclic ring; with the proviso that R_1 is not methoxy if m and n are both 1, R_2 is hydrogen and R_3 is fluoro; or a salt of the said compounds.

The general terms used hereinbefore and hereinafter preferably have within the context of this disclosure the following meanings, unless otherwise indicated:

Where the plural form is used for compounds, salts, and the like, this is taken to mean also a single compound, salt, or the like.

Asymmetric carbon atoms of a compound of formula I that are optionally present may exist in the (R), (S) or (R,S) configuration, preferably in the (R) or (S) configuration. Substituents at a double bond or a ring may be present in cis- (= Z-) or trans (= E-) form. The compounds may thus be present as mixtures of isomers or preferably as pure isomers.

Preferably alkyl contains up to 20 carbon atoms and is most preferably lower alkyl.

The prefix "lower" denotes a radical having up to and including a maximum of 7, especially up to and including a maximum of 4 carbon atoms, the radicals in question being either unbranched or branched with single or multiple branching.

Lower alkyl is, for example, methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec-butyl, tert-butyl, n-pentyl, isopentyl, neopentyl, n-hexyl or n-heptyl.

Lower alkyl R_2 is preferably methyl, ethyl or isopropyl.

Lower alkyl R_3 is preferably methyl or ethyl.

Substituted lower alkyl is lower alkyl as defined above where one or more substituents may be present, such as amino, N-lower alkylamino, N,N-di-lower alkylamino, N-lower alkanoylamino, N,N-di-lower alkanoylamino, hydroxy, lower alkoxy, lower alkanoyl, lower

alkanoyloxy, cyano, nitro, carboxy, lower alkoxy carbonyl, carbamoyl, amidino, guanidino, ureido, mercapto, lower alkylthio, halogen or a heterocyclic radical.

Substituted lower alkyl R_2 is preferably hydroxy-lower alkyl or amino-lower alkyl.

Mono- or di-substituted amino-sulfonyl is amino-sulfonyl, wherein the amino group is substituted by one or two radicals selected independently of one another from e.g. unsubstituted or substituted lower alkyl or a heterocyclic radical.

Unsubstituted, mono- or di-substituted amino-sulfonyl R_1 is preferably unsubstituted amino-sulfonyl.

Mono- or di-substituted amino is amino substituted by one or two radicals selected independently of one another from e.g. unsubstituted or substituted lower alkyl or a heterocyclic radical.

Mono- or di-substituted amino R_1 is preferably N-lower alkylamino or N,N-di-lower alkylamino, respectively.

Mono- or di-substituted amino R_4 is preferably N-lower alkylamino or N,N-di-lower alkylamino, respectively.

A heterocyclic radical contains especially up to 20 carbon atoms including possible substituents and is an unsaturated, partially unsaturated, or preferably saturated monocyclic radical having from 4 or 8 ring members and from 1 to 3 heteroatoms which are preferably selected from nitrogen, oxygen and sulfur, or a bi- or tri-cyclic radical wherein, for example, one or two benzene radicals are annellated (fused) to the mentioned monocyclic radical. The heterocyclic radical is optionally substituted by one or more radicals such as e.g. unsubstituted or substituted lower alkyl.

A heterocyclic radical R_1 is preferably lower alkyl-piperazinyl, especially 4-lower alkyl-piperazin-1-yl.

The heterocyclic ring formed by two vicinal R_1 substituents together with the phenyl carbon atoms to which they are attached contains especially up to 20 carbon atoms including possible substituents and is an unsaturated, partially unsaturated, or saturated monocyclic radical having from 4 or 8 ring members and from 1 to 3 heteroatoms which are preferably selected from nitrogen, oxygen and sulfur. The heterocyclic ring is optionally substituted by one or more radicals such as e.g. oxo (=O), thioxo (=S), or unsubstituted or substituted lower alkyl. Preferably this ring is a thiazol, 1-oxo-thiazol or dioxol ring.

Lower alkyl substituted by a heterocyclic radical R_1 is preferably lower alkyl substituted by lower alkyl-piperazinyl, especially by 4-lower alkyl-piperazin-1-yl.

A heterocyclic radical R_4 is preferably morpholinyl, especially morpholin-4-yl, or lower alkyl-piperidyl, especially 1-lower alkyl-piperidin-4-yl.

A heterocyclic radical R_5 is preferably lower alkyl-piperazinyl, especially 4-lower alkyl-piperazin-1-yl.

A heterocyclic radical R_2 is preferably piperidyl, such as piperidin-4-yl, lower alkyl-piperidyl, such as 1-lower alkyl-piperidin-4-yl, or tetrahydro-pyranyl, such as tetrahydro-pyran-4-yl.

Etherified hydroxy is, for example, alkoxy, especially lower alkoxy, such as methoxy, ethoxy and tert-butoxy.

Etherified hydroxy R_6 is preferably lower alkoxy, especially ethoxy or tert-butoxy.

Etherified hydroxy R_3 is preferably lower alkoxy, especially methoxy or ethoxy, or phenyl-lower alkoxy, wherein the phenyl moiety is preferably unsubstituted or substituted by e.g. unsubstituted or substituted lower alkyl.

Esterified hydroxy is preferably hydroxy esterified by an organic carboxylic acid, such as alcanoic acid, and is for example lower alkanoyloxy.

Halogen is primarily fluorine, chlorine, bromine, or iodine, especially fluorine, chlorine, or bromine.

Halogen R₃ is preferably chloro.

X is preferably -O-.

m is preferably 1 or 2.

n is preferably 1 or 2.

R₁ is preferably attached to the phenyl ring in the meta and/or para position, most preferably in the meta position.

R₃ is preferably attached to the phenyl ring in the meta and/or para position, most preferably in the para position.

R₄ is preferably mono- or di-substituted amino, or a heterocyclic radical.

Salts are especially the pharmaceutically acceptable salts of compounds of formula I.

Such salts are formed, for example, as acid addition salts, preferably with organic or inorganic acids, from compounds of formula I with a basic nitrogen atom, especially the pharmaceutically acceptable salts.

In the presence of negatively charged radicals, such as carboxy or sulfo, salts may also be formed with bases, e.g. metal or ammonium salts, such as alkali metal or alkaline earth metal salts, or ammonium salts with ammonia or suitable organic amines, such as tertiary monoamines.

In the presence of a basic group and an acid group in the same molecule, a compound of formula I may also form internal salts.

For isolation or purification purposes it is also possible to use pharmaceutically unacceptable salts, for example picrates or perchlorates. Only the pharmaceutically acceptable salts or

free compounds (if the occasion arises, in the form of pharmaceutical compositions) attain therapeutic use, and these are therefore preferred.

In view of the close relationship between the novel compounds in free form and in the form of their salts, including those salts that can be used as intermediates, for example in the purification or identification of the novel compounds, hereinbefore and hereinafter any reference to the free compounds is to be understood as referring also to the corresponding salts, as appropriate and expedient.

The compounds of formula I have valuable, pharmacologically useful properties. In particular they exhibit specific inhibitory activities that are of pharmacological interest. They are effective especially as protein tyrosine kinase inhibitors and/or (furthermore) as inhibitors of serine/threonine protein kinases; they exhibit, for example, powerful inhibition of the tyrosine kinase activity of the epidermal growth factor receptor (EGF-R) and of ErbB-2 Kinase. These two protein tyrosine kinase receptors, together with their family members ErbB-3 and ErbB-4, play a key role in signal transmission in a large number of mammalian cells, including human cells, especially epithelial cells, cells of the immune system and cells of the central and peripheral nervous system. For example, in various cell types, EGF-induced activation of receptor-associated protein tyrosine kinase is a prerequisite for cell division and hence for the proliferation of the cell population. Most importantly, overexpression of the EGF-R (HER-1) and/or ErbB-2 (HER-2) has been observed in substantial fractions of many human tumours. EGF-R, e.g., was found to be overexpressed in non small-cell lung cancers, squamous carcinoma (head and neck), breast, gastric, ovarian, colon and prostate cancers as well as in gliomas. ErbB-2 was found to be overexpressed in squamous carcinoma (head and neck), breast, gastric, and ovarian cancers as well as in gliomas.

In addition to inhibiting the tyrosine kinase activity of the EGF-R, the compounds of formula I also inhibit to varying extents other protein tyrosine kinases that are involved in signal transmission mediated by trophic factors, specially the vascular endothelial growth factor (VEGF) receptor family (e.g. KDR, Flt-1, Flt-3) but also abl kinase, especially v-abl, kinases from the family of Src, especially c-Src, Lck and Fyn, the other members of the EGF receptor family such as ErbB-3 (HER-3) and ErbB-4 (HER-4), CSF-1, Kit, FGF receptor and the cyclin-dependent kinases CDK1 and CDK2, all of

which play a part in growth regulation and transformation in mammalian cells, including human cells.

The inhibition of EGF-R tyrosine kinase activity can be demonstrated using known methods, for example using the recombinant intracellular domain of the EGF-receptor [EGF-R ICD; see, for example, E. McGlynn et al., *Europ. J. Biochem.* 207, 265-275 (1992)]. Compared with the control without inhibitor, the compounds of formula I inhibit the enzyme activity by 50 % (IC_{50}), for example in a concentration of from 0.0005 to 0.5 μ M, especially from 0.001 to 0.1 μ M.

As well as or instead of inhibiting EGF-R tyrosine kinase activity, the compounds of formula I also inhibit other members of this family of receptors, like ErbB-2. The inhibitory activity (IC_{50}) is approximately in the range of 0.001 to 0.5 μ M. The inhibition of ErbB-2 tyrosine kinase (HER-2) can be determined, for example, analogously to the method used for EGF-R protein tyrosine kinase [see C. House et al., *Europ. J. Biochem.* 140, 363-367 (1984)]. The ErbB-2 kinase can be isolated, and its activity determined, by means of protocols known *per se*, for example in accordance with T. Akiyama et al., *Science* 232, 1644 (1986).

Surprisingly, the compounds of formula I especially also inhibit the tyrosine kinase activity of the VEGF receptor family very potently. The compounds of the present invention are therefore very effective dual inhibitors of EGF- and VEGF-receptor family members. For inhibition of KDR and Flt-1 and inhibition of growth factor-induced proliferation of HUVECS see J. Wood et al., *Cancer Res.* 60, 2178-2189 (2000). The compounds of formula I inhibit e.g. the KDR tyrosine kinase activity with an IC_{50} of from about 1 nM to about 1 μ M, especially from about 5 nM to about 0.5 μ M.

The action of the compounds of formula I on EGF-induced phosphorylation of the EGF-R can be determined in the human A431 epithelial carcinoma cell line by means of an ELISA which is described in U. Trinks et al., *J. Med. Chem.* 37:7, 1015-1027 (1994). In that test (EGF-R ELISA) the compounds of formula I exhibit an IC_{50} of approximately from 0.001 to 1 μ M.

The compounds of formula I potently inhibit the growth of EGF-R overexpressing NCI-H596 non-small cell lung carcinoma cells [see e.g. W. Lei, et al., *Anticancer Res.* **19**(1A), 221-228 (1999)] at an IC_{50} of approximately 0.01 to 1 μ M. In the same range of activity, the compounds of formula I also potently inhibit the growth of ErbB-2-overexpressing BT474 human breast cancer cells. The test procedures are adapted from T. Meyer et al., *Int. J. Cancer* **43**, 851 (1989). The inhibitory activity of the compounds of formula I is determined, briefly, as follows: NCI-H596 cells (10 000/microtitre plate well) are transferred to 96-well microtitre plates. The test compounds [dissolved in dimethyl sulfoxide (DMSO)] are added in a series of concentrations (dilution series) in such a manner that the final concentration of DMSO is not greater than 1 % (v/v). After the addition, the plates are incubated for three days during which the control cultures without test compound are able to undergo at least three cell-division cycles. The growth of the NCI-H596 cells is measured by means of methylene blue staining: after the incubation the cells are fixed with glutaraldehyde, washed with water and stained with 0.05 % methylene blue. After a washing step the stain is eluted with 3 % HCl and the optical density (OD) per well of the microtitre plate is measured using a Titertek Multiskan (Titertek, Huntsville, AL, USA) at 665 nm. IC_{50} values are determined by a computer-aided system using the formula:

$$IC_{50} = [(OD_{test} - OD_{start}) / (OD_{control} - OD_{start})] \times 100.$$

The IC_{50} value in those experiments is given as that concentration of the test compound in question that results in a cell count that is 50 % lower than that obtained using the control without inhibitor. The compounds of formula I exhibit inhibitory activity with an IC_{50} in the range from approximately 0.01 to 1 μ M.

The compounds of formula I exhibit inhibition of the growth of tumour cells also *in vivo*, as shown, for example, by the test described below: the test is based on inhibition of the growth of the human squamous lung carcinoma cell line NCI-H596 [ATCC HTB 178; American Type Culture Collection, Rockville, Maryland, USA; see Santon, J.B., et al., *Cancer Research* **46**, 4701-4705 (1986) and Ozawa, S., et al., *Int. J. Cancer* **40**, 706-710 (1987)], which is transplanted into female BALB/c nude mice (Bomholtgard, Denmark). That carcinoma exhibits a growth that correlates with the extent of the expression of the EGF-R. Tumours are established after subcutaneous (s.c.) injection of cells [a minimum of 2×10^6 cells in 100 μ l phosphate-buffered saline (PBS) or medium] in carrier mice (4-8 mice). Injections are

made s.c. in the left flank of the mouse mid-way between the tail and the head. The resulting tumours are serially passaged for a minimum of three consecutive transplantations prior to start of the treatment. During this time tumour growth rates stabilize. Tumours are not passaged more than 12 times. For the therapy experiment tumour fragments of roughly 25 mg are transplanted s.c. into the left flank of the animals using a 13-gauge trocar needle under Forene® (Abbott, Switzerland) anaesthesia. Tumour growth and body weights are monitored twice per week. All treatments are initiated when the tumour attains a volume of 100 to 250 mm³. The tumour volumes are calculated using the known formula Length x Diameter² x $\pi/6$ [see Evans, B.D., et al., Brit. J. Cancer 45, 466-8 (1982)]. Antitumour activity is expressed as T/C % (mean increase of tumour volumes of treated animals divided by the mean increase of tumour volumes of control animals multiplied by 100 %). At a dose of from 3 to 100 mg/kg of active ingredient, distinct inhibition of the tumour growth is found, for example T/C % values of less than 50.

The compounds of formula I may inhibit other protein tyrosine kinases that are involved in signal transmission mediated by trophic factors, for example abl kinase, such as especially v-abl kinase (IC₅₀ for example from 0.01 to 5 μ M), kinases from the family of the src kinases, such as especially c-src kinase (IC₅₀ for example from 0.1 to 10 μ M) and serine/threonine kinases, for example protein kinase C, especially A-Raf, B-Raf and c-Raf and subfamilies thereof, all of these kinases being involved in growth regulation and transformation in mammalian cells, including human cells.

The above-mentioned inhibition of v-abl tyrosine kinase is determined by the methods of N. Lydon et al., Oncogene Research 5, 161-173 (1990) and J. F. Geissler et al., Cancer Research 52, 4492-4498 (1992). In those methods [Val⁵]-angiotensin II and [γ -³²P]-ATP are used as substrates.

The compounds of formula I which inhibit the tyrosine kinase activity of the EGF-R or of the other protein tyrosine kinases mentioned are therefore useful, for example, in the treatment of benign or malignant tumours. The compounds of formula I are e.g. able to simultaneously inhibit the growth of tumors with deregulated EGF-R and/or ErbB-2 activity as well as to inhibit the vascularisation of solid tumors triggered by VEGF. This combined activity leads to an improved antitumour effect (see also WO 02/41882). Moreover, the use of a dual inhibitor reduces the risk of drug-drug interactions and further reduces the total drug load as

compared to a combination therapy. The compounds of formula I are capable of slowing down tumor growth or effecting tumour regression and of preventing the formation of tumour metastases and the growth of micrometastases. They can be used especially in the case of epidermal hyperproliferation (psoriasis), in the treatment of neoplasias of epithelial character, e.g. mammary carcinomas, and in leukaemias. In addition, the compounds of formula I can be used in the treatment of those disorders of the immune system in which several or, especially, individual protein tyrosine kinases and/or (furthermore) serine/threonine protein kinases are involved; the compounds of formula I can also be used in the treatment of those disorders of the central or peripheral nervous system in which signal transmission by several or, especially, a single protein tyrosine kinase(s) and/or (furthermore) serine/threonine protein kinase(s) is/are involved.

In general, the present invention relates also to the use of the compounds of formula I for the inhibition of the mentioned protein kinases, in particular to their use for the dual inhibition of EGF- and VEGF-receptor family members.

The compounds according to the invention can be used both alone and in combination with other pharmacologically active compounds, for example together with inhibitors of the enzymes of polyamine synthesis, inhibitors of protein kinase C, inhibitors of other tyrosine kinases, cytokines, negative growth regulators, for example TGF- β or IFN- β , aromatase inhibitors, antioestrogens and/or cytostatic drugs.

With the groups of preferred compounds of formula I mentioned hereinafter, definitions of substituents from the general definitions mentioned hereinbefore may reasonably be used, for example, to replace more general definitions with more specific definitions or especially with definitions characterized as being preferred.

Preference is given to a compound of formula I, wherein

m and n are independently of one another an integer from 1 to 5;

R_1 is unsubstituted, mono- or di-substituted amino-sulfonyl; unsubstituted, mono- or di-substituted amino; a heterocyclic radical; lower alkyl substituted by a heterocyclic radical; a radical R_4 -lower alkyl-X-, wherein R_4 is hydrogen, unsubstituted, mono- or di-substituted amino, or a heterocyclic radical, and X is -S- or -O-; or a radical R_5 -C(=O)-, wherein R_5 is hydrogen, unsubstituted or substituted lower alkyl, free or etherified hydroxy, unsubstituted,

mono- or di-substituted amino, or a heterocyclic radical; wherein the R_1 substituents are selected independently of one another if $m > 1$;

or two vicinal R_1 substituents together with the phenyl carbon atoms to which they are attached form a heterocyclic ring;

R_2 is hydrogen; and

R_3 is halogen; unsubstituted or substituted lower alkyl; or free, etherified or esterified hydroxy; wherein the R_3 substituents are selected independently of one another if $n > 1$;

or two vicinal R_3 substituents together with the phenyl carbon atoms to which they are attached form a heterocyclic ring;

with the proviso that R_1 is not methoxy if m and n are both 1, R_2 is hydrogen and R_3 is fluoro; and the salts thereof.

Special preference is given to a compound of formula I, wherein

m and n are independently of one another an integer from 1 to 3;

R_1 is amino-sulfonyl; mono- or di-substituted amino; a heterocyclic radical; lower alkyl substituted by a heterocyclic radical; a radical R_4 -lower alkyl-X-, wherein R_4 is hydrogen, mono- or di-substituted amino, or a heterocyclic radical, and X is -S- or -O-; or a radical R_5 -C(=O)-, wherein R_5 is lower alkyl, free or etherified hydroxy, or a heterocyclic radical;

wherein the R_1 substituents are selected independently of one another if $m > 1$;

or two vicinal R_1 substituents together with the phenyl carbon atoms to which they are attached form a heterocyclic ring;

R_2 is hydrogen; and

R_3 is halogen; lower alkyl; or free or etherified hydroxy; wherein the R_3 substituents are selected independently of one another if $n > 1$;

or two vicinal R_3 substituents together with the phenyl carbon atoms to which they are attached form a heterocyclic ring;

with the proviso that R_1 is not methoxy if m and n are both 1, R_2 is hydrogen and R_3 is fluoro; and the salts thereof.

Very special preference is given to a compound of formula I, wherein

m is an integer from 1 to 3 and n is 1 or 2;

R_1 is amino-sulfonyl; N,N-di-lower alkylamino; lower alkyl-piperazinyl; lower alkyl substituted by lower alkyl-piperazinyl; a radical R_4 -lower alkyl-X-, wherein R_4 is hydrogen, N,N-di-lower alkylamino, morpholinyl or lower alkyl-piperidyl, and X is -S- or -O-; or a radical R_5 -C(=O)-,

wherein R_5 is lower alkyl, hydroxy, lower alkoxy or lower alkyl-piperazinyl; wherein the R_1 substituents are selected independently of one another if $m > 1$;
or two vicinal R_1 substituents together with the phenyl carbon atoms to which they are attached form a thiazol or 1-oxo-thiazol ring;
 R_2 is hydrogen; and
 R_3 is halogen, lower alkyl, hydroxy, lower alkoxy or phenyl-lower alkoxy; wherein the R_3 substituents are selected independently of one another if n is 2;
or two R_3 radicals together form a dioxol ring;
with the proviso that R_1 is not methoxy if m and n are both 1, R_2 is hydrogen and R_3 is fluoro;
and the salts thereof.

Preference is further given to a compound of formula I, wherein
 m and n are independently of one another an integer from 1 to 5;
 R_1 is unsubstituted, mono- or di-substituted amino-sulfonyl; unsubstituted, mono- or di-substituted amino; a heterocyclic radical; lower alkyl substituted by a heterocyclic radical; or a radical $R_5-C(=O)-$, wherein R_5 is hydrogen, unsubstituted or substituted lower alkyl, free or etherified hydroxy, unsubstituted, mono- or di-substituted amino, or a heterocyclic radical;
wherein the R_1 substituents are selected independently of one another if $m > 1$;
or two vicinal R_1 substituents together with the phenyl carbon atoms to which they are attached form a heterocyclic ring;
 R_2 is hydrogen, unsubstituted or substituted lower alkyl or a heterocyclic radical; and
 R_3 is halogen; unsubstituted or substituted lower alkyl; or free, etherified or esterified hydroxy; wherein the R_3 substituents are selected independently of one another if $n > 1$;
or two vicinal R_3 substituents together with the phenyl carbon atoms to which they are attached form a heterocyclic ring;
and the salts thereof.

Preference is further also given to a compound of formula I, wherein
 m and n are independently of one another an integer from 1 to 5;
 R_1 is unsubstituted, mono- or di-substituted amino-sulfonyl; unsubstituted, mono- or di-substituted amino; a heterocyclic radical; lower alkyl substituted by a heterocyclic radical; or a radical $R_5-C(=O)-$, wherein R_5 is hydrogen, unsubstituted or substituted lower alkyl, free or etherified hydroxy, unsubstituted, mono- or di-substituted amino, or a heterocyclic radical;
wherein the R_1 substituents are selected independently of one another if $m > 1$;

or two vicinal R₁ substituents together with the phenyl carbon atoms to which they are attached form a heterocyclic ring;

R₂ is hydrogen; and

R₃ is halogen; unsubstituted or substituted lower alkyl; or free, etherified or esterified hydroxy; wherein the R₃ substituents are selected independently of one another if n>1; or two vicinal R₃ substituents together with the phenyl carbon atoms to which they are attached form a heterocyclic ring;

and the salts thereof.

Special preference is further given to a compound of formula I, wherein m and n are independently of one another an integer from 1 to 3;

R₁ is amino-sulfonyl; mono- or di-substituted amino; a heterocyclic radical; lower alkyl substituted by a heterocyclic radical; or a radical R₅-C(=O)-, wherein R₅ is lower alkyl, free or etherified hydroxy, or a heterocyclic radical; wherein the R₁ substituents are selected independently of one another if m>1;

or two vicinal R₁ substituents together with the phenyl carbon atoms to which they are attached form a heterocyclic ring;

R₂ is hydrogen; and

R₃ is halogen; lower alkyl; or free or etherified hydroxy; wherein the R₃ substituents are selected independently of one another if n>1;

or two vicinal R₃ substituents together with the phenyl carbon atoms to which they are attached form a heterocyclic ring;

and the salts thereof.

Very special preference is further given to a compound of formula I, wherein

m and n are independently of one another 1 or 2;

R₁ is amino-sulfonyl; N,N-di-lower alkylamino; lower alkyl-piperazinyl; lower alkyl substituted by lower alkyl-piperazinyl; or a radical R₅-C(=O)-, wherein R₅ is lower alkyl, hydroxy, lower alkoxy or lower alkyl-piperazinyl; wherein the R₁ substituents are selected independently of one another if m is 2;

or two vicinal R₁ substituents together with the phenyl carbon atoms to which they are attached form a thiazol or 1-oxo-thiazol ring;

R₂ is hydrogen; and

R_3 is halogen, lower alkyl, hydroxy, or lower alkoxy; wherein the R_3 substituents are selected independently of one another if n is 2; and the salts thereof.

Very special preference is also given to a compound of formula I, wherein R_1 is a heterocyclic radical, and the salts thereof.

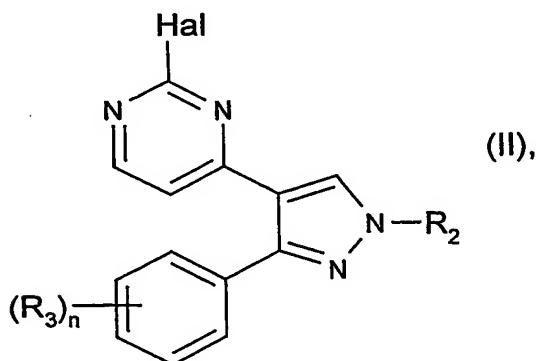
Very special preference is further given to a compound of formula I mentioned in the Examples below, or a salt, especially a pharmaceutically acceptable salt, thereof.

Also especially preferred are compounds of formula I, which - according to the above-described tyrosine kinase inhibition assays - inhibit HER-1, HER-2 and KDR with IC_{50} values of less than 300 nM, most preferably of less than 100 nM.

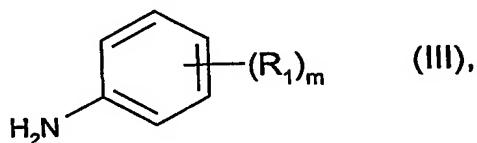
Very special preference is further given to compounds of formula I which inhibit the tyrosine kinase activity of at least one member of the EGF receptor family together with at least one member of the VEGF receptor family (dual inhibition of EGF- and VEGF-receptor family members) with IC_{50} values in the range of 0.5 nM to 0.5 μ M, especially in the range of 1 nM to 300 nM, based on the above-described tyrosine kinase inhibition assays.

The compounds of formula I or salts thereof are prepared in accordance with processes known per se, though not previously described for the manufacture of the compounds of the formula I, especially whereby

- a) in order to prepare a compound of formula I, wherein R_2 is hydrogen or lower alkyl,
 - (i) a compound of formula II

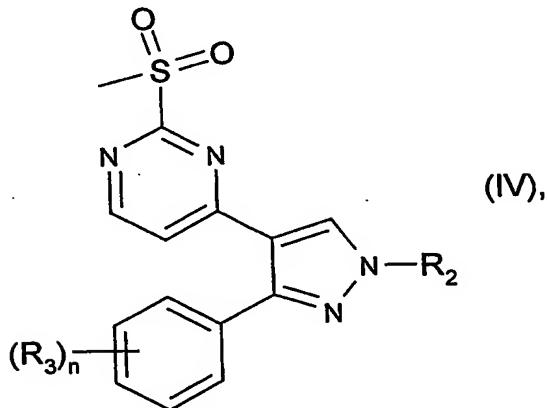


wherein Hal is halogen, R₂ is hydrogen or lower alkyl and n and R₃ have the meanings as defined for a compound of formula I, is reacted with a compound of formula III



wherein m and R₁ have the meanings as defined for a compound of formula I, or

(ii) a compound of formula IV



wherein R₂ is hydrogen or lower alkyl and n and R₃ have the meanings as defined for a compound of formula I, is reacted with a compound of formula III; or

b) in order to prepare a compound of formula I, wherein R₂ is unsubstituted or substituted lower alkyl or a heterocyclic radical, a compound of formula I, wherein R₂ is hydrogen, is reacted with a compound of the formula R₂-OH, wherein R₂ is unsubstituted or substituted lower alkyl or a heterocyclic radical wherein the unsubstituted or substituted lower alkyl or the heterocyclic radical is attached to the hydroxy group of R₂-OH via a carbon atom of the lower alkyl moiety or via a ring carbon atom of the heterocyclic radical, respectively;

whereby functional groups which are present in the starting compounds of process a) or b) and are not intended to take part in the reaction, are present in protected form if necessary,

and protecting groups that are present are cleaved, whereby the said starting compounds may also exist in the form of salts provided that a salt-forming group is present and a reaction in salt form is possible;

and, if so desired, a compound of formula I thus obtained is converted into another compound of formula I, a free compound of formula I is converted into a salt, an obtained salt of a compound of formula I is converted into the free compound or another salt, and/or a mixture of isomeric compounds of formula I is separated into the individual isomers.

Description of the process variants:

Regarding process a):

- (i) The reaction between a compound of formula II and a compound of formula III preferably takes place in a suitable inert solvent such as dioxane, in the presence of an acid such as HCl, at elevated temperature, preferably at around 100 °C. In a compound of formula II, Hal is preferably chloro or bromo, especially bromo.
- (ii) The reaction between a compound of formula IV and a compound of formula III preferably takes place under those conditions described for the analogous procedure in Klutchko et al., Journal of Medicinal Chemistry, 1998, Vol. 41, No. 17, 3276-3292.

Regarding process b):

The reaction between a compound of formula I, wherein R₂ is hydrogen, and a compound of the formula R₂-OH preferably takes place under the Mitsunobu reaction conditions such as those described in: Mitsunobu, Oyo; Synthesis 1981, p. 1 - 27.

Additional process steps

In the additional process steps, carried out as desired, functional groups of the starting compounds which should not take part in the reaction may be present in unprotected form or may be protected for example by one or more protecting groups. The protecting groups are then wholly or partly removed according to one of the known methods.

Protecting groups, and the manner in which they are introduced and removed are described, for example, in "Protective Groups in Organic Chemistry", Plenum Press, London, New York 1973, and in "Methoden der organischen Chemie", Houben-Weyl, 4th edition, Vol. 15/1, Georg-Thieme-Verlag, Stuttgart 1974 and in Theodora W. Greene, "Protective Groups in

"Organic Synthesis", John Wiley & Sons, New York 1981. A characteristic of protecting groups is that they can be removed readily, i.e. without the occurrence of undesired secondary reactions, for example by solvolysis, reduction, photolysis or alternatively under physiological conditions.

The end products of formula I may however also contain substituents that can also be used as protecting groups in starting materials for the preparation of other end products of formula I. Thus, within the scope of this text, only a readily removable group that is not a constituent of the particular desired end product of formula I is designated a "protecting group", unless the context indicates otherwise.

General process conditions

All process steps described here can be carried out under known reaction conditions, preferably under those specifically mentioned, in the absence of or usually in the presence of solvents or diluents, preferably those that are inert to the reagents used and able to dissolve them, in the absence or presence of catalysts, condensing agents or neutralising agents, for example ion exchangers, typically cation exchangers, for example in the H⁺ form, depending on the type of reaction and/or reactants at reduced, normal, or elevated temperature, for example in the range from -100 °C to about 190 °C, preferably from about -80 °C to about 150 °C, for example at -80 to -60 °C, at RT, at -20 to 40 °C, at 0 to 100 °C or at the boiling point of the solvent used, under atmospheric pressure or in a closed vessel, if need be under pressure, and/or in an inert, for example an argon or nitrogen, atmosphere.

The invention relates also to those embodiments of the process in which one starts from a compound obtainable at any stage as an intermediate and carries out the missing steps, or breaks off the process at any stage, or forms a starting material under the reaction conditions, or uses said starting material in the form of a reactive derivative or salt, or produces a compound obtainable by means of the process according to the invention under those process conditions, and further processes the said compound *in situ*. In the preferred embodiment, one starts from those starting materials which lead to the compounds described hereinabove as preferred.

In the preferred embodiment, a compound of formula I is prepared according to the processes and process steps defined in the Examples.

The compounds of formula I, including their salts, are also obtainable in the form of hydrates, or their crystals can include for example the solvent used for crystallisation (present as solvates).

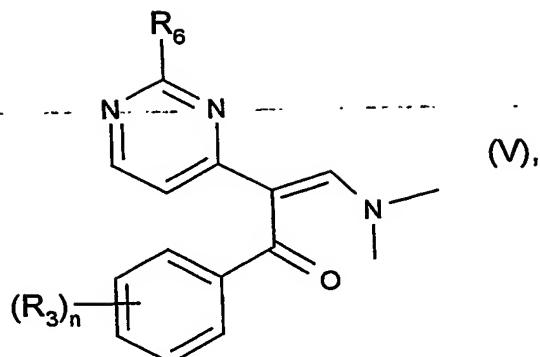
Starting materials

New starting materials and/or intermediates, as well as processes for the preparation thereof, are likewise the subject of this invention. In the preferred embodiment, such starting materials are used and reaction conditions so selected as to enable the preferred compounds to be obtained.

The starting materials used in the above described processes a) to b) are known, capable of being prepared according to known processes, or commercially obtainable; in particular, they can be prepared using processes as described in the Examples.

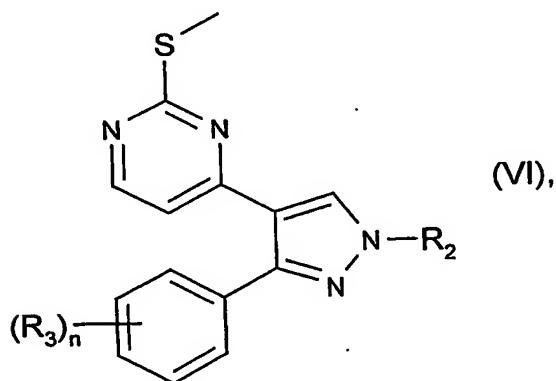
In the preparation of starting materials, existing functional groups which do not participate in the reaction should, if necessary, be protected. Preferred protecting groups, their introduction and their removal are described above or in the Examples. In place of the respective starting materials and transients, salts thereof may also be used for the reaction, provided that salt-forming groups are present and the reaction with a salt is also possible. Where the term starting materials is used hereinbefore and hereinafter, the salts thereof are always included, insofar as reasonable and possible.

A compound of formula II, wherein R₂ is hydrogen or lower alkyl and n and R₃ have the meanings as defined for a compound of formula I, can be prepared for example by reacting a compound of formula V



wherein R_6 is halogen and n and R_3 have the meanings as defined for a compound of formula I, with hydrazine (H_2N-NH_2) or N-lower alkyl-hydrazine (lower alkyl- $NH-NH_2$), respectively, in a suitable solvent, e.g. lower alcohols, such as ethanol, preferably at around room temperature.

A compound of formula IV, wherein R_2 is hydrogen or lower alkyl and n and R_3 have the meanings as defined for a compound of formula I, can be prepared for example by reacting a compound of formula VI

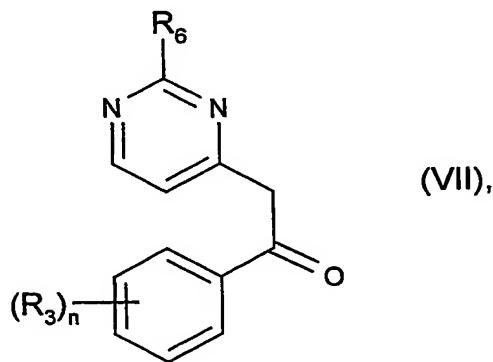


wherein R_2 is hydrogen or lower alkyl and n and R_3 have the meanings as defined for a compound of formula I, with 3-chloroperoxybenzoic acid in $CHCl_3$, e.g. under those conditions described for the analogous procedure in Klutchko et al., Journal of Medicinal Chemistry, 1998, Vol. 41, No. 17, 3276-3292.

A compound of formula VI, wherein R_2 is hydrogen or lower alkyl and n and R_3 have the meanings as defined for a compound of formula I, can be prepared for example by reacting a compound of formula V, wherein R_6 is $-S-CH_3$ and n and R_3 have the meanings as defined

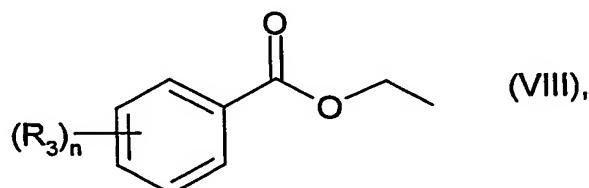
for a compound of formula I, with hydrazine ($\text{H}_2\text{N-NH}_2$) or N-lower alkyl-hydrazine (lower alkyl- NH-NH_2), respectively, in a suitable solvent, e.g. lower alcohols, such as ethanol, preferably at around room temperature.

A compound of formula V, wherein R_6 is halogen or $-\text{S-CH}_3$ and n and R_3 have the meanings as defined for a compound of formula I, can be prepared for example by reacting a compound of formula VII

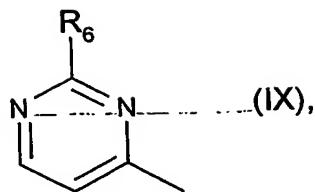


wherein R_6 is halogen or $-\text{S-CH}_3$, respectively, and n and R_3 have the meanings as defined for a compound of formula I, with $\text{N,N-dimethylformamid-dimethylacetal}$, at elevated temperature, preferably at around $100\text{ }^\circ\text{C}$.

A compound of formula VII, wherein R_6 is halogen and n and R_3 have the meanings as defined for a compound of formula I, can be prepared for example by reacting a compound of formula VIII

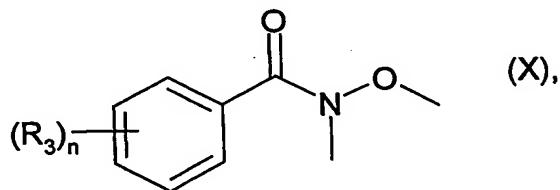


wherein n and R_3 have the meanings as defined for a compound of formula I, with a compound of formula IX



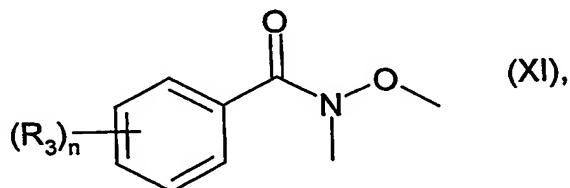
wherein R_6 is halogen, in the presence of lithiumdiisopropylamide, in a suitable organic solvent or mixture of solvents, preferably starting the reaction at reduced temperature, preferably at around -75°C , and letting it to reach room temperature.

A compound of formula VII, wherein R_6 is $-S-\text{CH}_3$ and n and R_3 have the meanings as defined for a compound of formula I, can be prepared for example by reacting a compound of formula X



wherein n and R_3 have the meanings as defined for a compound of formula I, with a compound of formula IX, wherein R_6 is $-S-\text{CH}_3$, in the presence of lithiumdiisopropylamide, in a suitable organic solvent or mixture of solvents, preferably starting the reaction at reduced temperature, preferably at around -75°C , and letting it to reach room temperature.

A compound of formula X, wherein n and R_3 have the meanings as defined for a compound of formula I, can be prepared for example by reacting a compound of formula XI



wherein n and R_3 have the meanings as defined for a compound of formula I, with N-O-dimethylhydroxylamine HCl in CH_2Cl_2 , e.g. under those conditions described for the analogous procedure in Nahm, Steven; Weinreb, Steven M.; *Tetrahedron Lett.*; 1981; 22 (39); 3815-3818.

The remaining starting materials are known, capable of being prepared according to known processes, or commercially available; or in particular, they can be prepared using processes as described in the Examples.

Pharmaceutical compositions, methods, and uses

The present invention relates also to pharmaceutical compositions that comprise a compound of formula I, or a pharmaceutically acceptable salt thereof, as active ingredient and that can be used especially in the treatment of the diseases mentioned at the beginning. Compositions for enteral administration, such as nasal, buccal, rectal or, especially, oral administration, and for parenteral administration, such as intravenous, intramuscular or subcutaneous administration, to warm-blooded animals, especially humans, are especially preferred. The compositions contain the active ingredient alone or, preferably, together with a pharmaceutically acceptable carrier. The dosage of the active ingredient depends upon the disease to be treated and upon the species, its age, weight, and individual condition, the individual pharmacokinetic data, and the mode of administration.

The present invention also relates to pro-drugs of a compound of formula I that convert *in vivo* to the compound of formula I as such. Any reference to a compound of formula I is therefore to be understood as referring also to the corresponding pro-drugs of the compound of formula I, as appropriate and expedient.

The invention relates also to compounds of formula I, or a pharmaceutically acceptable salt thereof, as such or in the form of a pharmaceutical composition, for use in a method for the prophylactic or especially therapeutic treatment of the human or animal body, to a process for the preparation thereof (especially in the form of compositions for the treatment of tumours) and to a method of treating proliferative diseases, primarily tumour diseases, especially those mentioned above.

The invention relates also to processes and to the use of compounds of formula I, or a pharmaceutically acceptable salt thereof, for the preparation of pharmaceutical compositions which comprise compounds of formula I, or a pharmaceutically acceptable salt thereof, as active component (active ingredient).

If desired, the said pharmaceutical compositions may also contain further active components, for example cytostatics, and/or may be used in combination with known therapeutic processes, for example the administration of hormones or radiation.

Preference is given for a pharmaceutical composition which is suitable for administration to a warm-blooded animal, especially humans or commercially useful mammals suffering from a disease which responds to an inhibition of a protein tyrosine kinase, especially to a dual inhibition of EGF- and VEGF-receptor family members, especially a neoplastic disease, comprising an effective quantity of a compound of formula I for the inhibition of a protein tyrosine kinase, especially for the dual inhibition of EGF- and VEGF-receptor family members, or a pharmaceutically acceptable salt thereof, together with at least one pharmaceutically acceptable carrier.

A pharmaceutical composition for the prophylactic or especially therapeutic management of neoplastic and other proliferative diseases of a warm-blooded animal, especially a human or a commercially useful mammal requiring such treatment, especially suffering from such a disease, comprising as active ingredient in a quantity that is prophylactically or especially therapeutically active against said diseases a compound of formula I, or a pharmaceutically acceptable salt thereof, is likewise preferred.

The pharmaceutical compositions comprise from approximately 1% to approximately 95% active ingredient, single-dose administration forms comprising in the preferred embodiment from approximately 20% to approximately 90% active ingredient and forms that are not of single-dose type comprising in the preferred embodiment from approximately 5% to approximately 20% active ingredient. Unit dose forms are, for example, coated and uncoated tablets, ampoules, vials, suppositories or capsules. Examples are capsules containing from about 0.05 g to about 1.0 g of active substance.

The pharmaceutical compositions of the present invention are prepared in a manner known *per se*, for example by means of conventional mixing, granulating, coating, dissolving or lyophilising processes.

The invention relates likewise to a process or a method for the treatment of one of the pathological conditions mentioned hereinabove, especially a disease which responds to an

inhibition of a protein tyrosine kinase, especially to a dual inhibition of EGF- and VEGF-receptor family members, especially a corresponding neoplastic disease. The compounds of formula I, or pharmaceutically acceptable salts thereof, can be administered as such or in the form of pharmaceutical compositions, prophylactically or therapeutically, preferably in an amount effective against the said diseases, to a warm-blooded animal, for example a human, requiring such treatment, the compounds especially being used in the form of pharmaceutical compositions. In the case of an individual having a bodyweight of about 70 kg the daily dose administered is from approximately 0.1 g to approximately 5 g, preferably from approximately 0.5 g to approximately 2 g, of a compound of the present invention.

The present invention relates especially also to the use of a compound of formula I, or a pharmaceutically acceptable salt thereof, especially a compound of formula I which is said to be preferred, or a pharmaceutically acceptable salt thereof, as such or in the form of a pharmaceutical composition with at least one pharmaceutically acceptable carrier, for the therapeutic and also prophylactic management of one or more of the diseases mentioned hereinabove, preferably a disease which responds to an inhibition of a protein tyrosine kinase, especially to a dual inhibition of EGF- and VEGF-receptor family members, especially a neoplastic disease, in particular if the said disease responds to an inhibition of a protein tyrosine kinase, especially to a dual inhibition of EGF- and VEGF-receptor family members.

The present invention relates especially also to the use of a compound of formula I, or a pharmaceutically acceptable salt thereof, especially a compound of formula I which is said to be preferred, or a pharmaceutically acceptable salt thereof, for the preparation of a pharmaceutical composition for the therapeutic and also prophylactic management of one or more of the diseases mentioned hereinabove, especially a neoplastic disease, in particular if the disease responds to an inhibition of a protein tyrosine kinase, especially to a dual inhibition of EGF- and VEGF-receptor family members.

A compound of formula I may also be used to advantage in combination with other antiproliferative agents. Such antiproliferative agents include, but are not limited to aromatase inhibitors, antiestrogens, topoisomerase I inhibitors, topoisomerase II inhibitors, microtubule active agents, alkylating agents, histone deacetylase inhibitors, farnesyl transferase inhibitors, COX-2 inhibitors, MMP inhibitors, mTOR inhibitors, antineoplastic

antimetabolites, platin compounds, compounds decreasing the protein kinase activity and further anti-angiogenic compounds, gonadorelin agonists, anti-androgens, bengamides, bisphosphonates, antiproliferative antibodies and temozolomide (TEMODAL®).

The term "aromatase inhibitors" as used herein relates to compounds which inhibit the estrogen production, i.e. the conversion of the substrates androstanedione and testosterone to estrone and estradiol, respectively. The term includes, but is not limited to steroids, especially exemestane and formestane and, in particular, non-steroids, especially aminoglutethimide, vorozole, fadrozole, anastrozole and, very especially, letrozole. Exemestane can be administered, e.g., in the form as it is marketed, e.g. under the trademark AROMASIN™. Formestane can be administered, e.g., in the form as it is marketed, e.g. under the trademark LENTARON™. Fadrozole can be administered, e.g., in the form as it is marketed, e.g. under the trademark AFEMA™. Anastrozole can be administered, e.g., in the form as it is marketed, e.g. under the trademark ARIMIDEX™. Letrozole can be administered, e.g., in the form as it is marketed, e.g. under the trademark FEMARA™ or FEMAR™. Aminoglutethimide can be administered, e.g., in the form as it is marketed, e.g. under the trademark ORIMETEN™.

A combination of the invention comprising an antineoplastic agent which is an aromatase inhibitor is particularly useful for the treatment of hormone receptor positive breast tumors.

The term "antiestrogens" as used herein relates to compounds which antagonize the effect of estrogens at the estrogen receptor level. The term includes, but is not limited to tamoxifen, fulvestrant, raloxifene and raloxifene hydrochloride. Tamoxifen can be administered, e.g., in the form as it is marketed, e.g. under the trademark NOLVADEX™. Raloxifene hydrochloride can be administered, e.g., in the form as it is marketed, e.g. under the trademark EVISTA™. Fulvestrant can be formulated as disclosed in US 4,659,516 or it can be administered, e.g., in the form as it is marketed, e.g. under the trademark FASLODEX™.

The term "topoisomerase I inhibitors" as used herein includes, but is not limited to topotecan, irinotecan, 9-nitrocamptothecin and the macromolecular camptothecin conjugate PNU-166148 (compound A1 in WO99/17804). Irinotecan can be administered, e.g., in the form as it is marketed, e.g. under the trademark CAMPTOSAR™. Topotecan can be administered, e.g., in the form as it is marketed, e.g. under the trademark HYCAMTIN™.

The term "topoisomerase II inhibitors" as used herein includes, but is not limited to the antracyclines doxorubicin (including liposomal formulation, e.g. CAELYX™), epirubicin, idarubicin and nemorubicin, the anthraquinones mitoxantrone and losoxantrone, and the podophyllotoxines etoposide and teniposide. Etoposide can be administered, e.g., in the form as it is marketed, e.g. under the trademark ETOPOPHOS™. Teniposide can be administered, e.g., in the form as it is marketed, e.g. under the trademark VM 26-BRISTOL™. Doxorubicin can be administered, e.g., in the form as it is marketed, e.g. under the trademark ADRIBLASTIN™. Epirubicin can be administered, e.g., in the form as it is marketed, e.g. under the trademark FARMORUBICIN™. Idarubicin can be administered, e.g., in the form as it is marketed, e.g. under the trademark ZAVEDOS™. Mitoxantrone can be administered, e.g., in the form as it is marketed, e.g. under the trademark NOVANTRON™.

The term "microtubule active agents" relates to microtubule stabilizing and microtubule destabilizing agents including, but not limited to the taxanes paclitaxel and docetaxel, the vinca alkaloids, e.g., vinblastine, especially vinblastine sulfate, vincristine especially vincristine sulfate, and vinorelbine, discodermolide and epothilones, such as epothilone B and D. Docetaxel can be administered, e.g., in the form as it is marketed, e.g. under the trademark TAXOTERE™. Vinblastine sulfate can be administered, e.g., in the form as it is marketed, e.g. under the trademark VINBLASTIN R.P.™. Vincristine sulfate can be administered, e.g., in the form as it is marketed, e.g. under the trademark FARMISTINTM. Discodermolide can be obtained, e.g., as disclosed in US 5,010,099.

The term "alkylating agents" as used herein includes, but is not limited to cyclophosphamide, ifosfamide and melphalan. Cyclophosphamide can be administered, e.g., in the form as it is marketed, e.g. under the trademark CYCLOSTIN™. Ifosfamide can be administered, e.g., in the form as it is marketed, e.g. under the trademark HOLOXAN™.

The term "histone deacetylase inhibitors" relates to compounds which inhibit the histone deacetylase and which possess antiproliferative activity.

The term "farnesyl transferase inhibitors" relates to compounds which inhibit the farnesyl transferase and which possess antiproliferative activity.

The term "COX-2 inhibitors" relates to compounds which inhibit the cyclooxygenase type 2 enzyme (COX-2) and which possess antiproliferative activity such as celecoxib (Celebrex®), rofecoxib (Vioxx®) and lumiracoxib (COX189):

The term "MMP inhibitors" relates to compounds which inhibit the matrix metalloproteinase (MMP) and which possess antiproliferative activity.

The term "mTOR inhibitors" relates to compounds which inhibit the mammalian target of rapamycin (mTOR) and which possess antiproliferative activity such as sirolimus (Rapamune®), everolimus (Certican™), CCI-779 and ABT578.

The term "antineoplastic antimetabolites" includes, but is not limited to 5-fluorouracil, tegafur, capecitabine, cladribine, cytarabine, fludarabine phosphate, fluorouridine, gemcitabine, 6-mercaptopurine, hydroxyurea, methotrexate, edatrexate and salts of such compounds, and furthermore ZD 1694 (RALTITREXED™), LY231514 (ALIMTA™), LY264618 (LOMOTREXOL™) and OGT719.

The term "platin compounds" as used herein includes, but is not limited to carboplatin, cis-platin and oxaliplatin. Carboplatin can be administered, e.g., in the form as it is marketed, e.g. under the trademark CARBOPLAT™. Oxaliplatin can be administered, e.g., in the form as it is marketed, e.g. under the trademark ELOXATIN™.

The term "compounds decreasing the protein kinase activity and further anti-angiogenic compounds" as used herein includes, but is not limited to compounds which decrease the activity of e.g. the Vascular Endothelial Growth Factor (VEGF), the Epidermal Growth Factor (EGF), c-Src, protein kinase C, Platelet-derived Growth Factor (PDGF), Bcr-Abl tyrosine kinase, c-kit, Flt-3 and Insulin-like Growth Factor I Receptor (IGF-IR) and Cyclin-dependent kinases (CDKs), and anti-angiogenic compounds having another mechanism of action than decreasing the protein kinase activity.

Compounds which decrease the activity of VEGF are especially compounds which inhibit the VEGF receptor, especially the tyrosine kinase activity of the VEGF receptor, and compounds binding to VEGF, and are in particular those compounds, proteins and monoclonal antibodies generically and specifically disclosed in WO 98/35958 (describing compounds of formula I), WO 00/09495, WO 00/27820, WO 00/59509, WO 98/11223, WO 00/27819, WO

01/55114, WO 01/58899 and EP 0 769 947; those as described by M. Prewett et al in Cancer Research 59 (1999) 5209-5218, by F. Yuan et al in Proc. Natl. Acad. Sci. USA, vol. 93, pp. 14765-14770, December 1996, by Z. Zhu et al in Cancer Res. 58, 1998, 3209-3214, and by J. Mordenti et al in Toxicologic Pathology, vol. 27, no. 1, pp 14-21, 1999; in WO 00/37502 and WO 94/10202; AngiostatinTM, described by M. S. O'Reilly et al, Cell 79, 1994, 315-328; and EndostatinTM, described by M. S. O'Reilly et al, Cell 88, 1997, 277-285; compounds which decrease the activity of EGF are especially compounds which inhibit the EGF receptor, especially the tyrosine kinase activity of the EGF receptor, and compounds binding to EGF, and are in particular those compounds generically and specifically disclosed in WO 97/02266 (describing compounds of formula IV), EP 0 564 409, WO 99/03854, EP 0520722, EP 0 566 226, EP 0 787 722, EP 0 837 063, WO 98/10767, WO 97/30034, WO 97/49688, WO 97/38983 and, especially, WO 96/33980; compounds which decrease the activity of c-Src include, but are not limited to, compounds inhibiting the c-Src protein tyrosine kinase activity as defined below and to SH2 interaction inhibitors such as those disclosed in WO97/07131 and WO97/08193; compounds inhibiting the c-Src protein tyrosine kinase activity include, but are not limited to, compounds belonging to the structure classes of pyrrolopyrimidines, especially pyrrolo[2,3-d]pyrimidines, purines, pyrazopyrimidines, especially pyrazo[3,4-d]pyrimidines, pyrazopyrimidines, especially pyrazo[3,4-d]pyrimidines and pyridopyrimidines, especially pyrido[2,3-d]pyrimidines. Preferably, the term relates to those compounds disclosed in WO 96/10028, WO 97/28161, WO97/32879 and WO97/49706; compounds which decreases the activity of the protein kinase C are especially those staurosporine derivatives disclosed in EP 0 296 110 (pharmaceutical preparation described in WO 00/48571) which compounds are protein kinase C inhibitors; further specific compounds that decrease protein kinase activity and which may also be used in combination with the compounds of the present invention are Imatinib (Gleevec®/Glivec®), PKC412, IressaTM (ZD1839), PKI166, PTK787, ZD6474, GW2016, CHIR-200131, CEP-7055/CEP-5214, CP-547632 and KRN-633; anti-angiogenic compounds having another mechanism of action than decreasing the protein kinase activity include, but are not limited to e.g. thalidomide (THALOMID), celecoxib (Celebrex), SU5416 and ZD6126.

The term "gonadorelin agonist" as used herein includes, but is not limited to abarelix, goserelin and goserelin acetate. Goserelin is disclosed in US 4,100,274 and can be administered, e.g., in the form as it is marketed, e.g. under the trademark ZOLADEX™. Abarelix can be formulated, e.g. as disclosed in US 5,843,901.

The term "anti-androgens" as used herein includes, but is not limited to bicalutamide (CASODEX™), which can be formulated, e.g. as disclosed in US 4,636,505.

The term "bengamides" relates to bengamides and derivatives thereof having aniproliferative properties.

The term "bisphosphonates" as used herein includes, but is not limited to etridonic acid, clodronic acid, tiludronic acid, pamidronic acid, alendronic acid, ibandronic acid, risedronic acid and zoledronic acid. "Etridonic acid" can be administered, e.g., in the form as it is marketed, e.g. under the trademark DIDRONEL™. "Clodronic acid" can be administered, e.g., in the form as it is marketed, e.g. under the trademark BONEFOS™. "Tiludronic acid" can be administered, e.g., in the form as it is marketed, e.g. under the trademark SKELID™. "Pamidronic acid" can be administered, e.g., in the form as it is marketed, e.g. under the trademark AREDIA™. "Alendronic acid" can be administered, e.g., in the form as it is marketed, e.g. under the trademark FOSAMAX™. "Ibandronic acid" can be administered, e.g., in the form as it is marketed, e.g. under the trademark BONDTRANAT™. "Risedronic acid" can be administered, e.g., in the form as it is marketed, e.g. under the trademark ACTONEL™. "Zoledronic acid" can be administered, e.g., in the form as it is marketed, e.g. under the trademark ZOMETA™.

The term "antiproliferative antibodies" as used herein includes, but is not limited to trastuzumab (Herceptin™), Trastuzumab-DM1, erlotinib (Tarceva™), bevacizumab (Avastin™), rituximab (Rituxan®), PRO64553 (anti-CD40) and 2C4 Antibody.

For the treatment of AML, compounds of formula I can be used in combination with standard leukemia therapies, especially in combination with therapies used for the treatment of AML. In particular, compounds of formula I can be administered in combination with e.g. farnesyltransferase inhibitors and/or other drugs used for the treatment of AML, such as

Daunorubicin, Adriamycin, Ara-C, VP-16, Teniposide, Mitoxantrone, Idarubicin, Carboplatinum and PKC412.

The structure of the active agents identified by code nos., generic or trade names may be taken from the actual edition of the standard compendium "The Merck Index" or from databases, e.g. Patents International (e.g. IMS World Publications).

The above-mentioned compounds, which can be used in combination with a compound of formula I, can be prepared and administered as described in the art such as in the documents cited above.

Examples:

The following Examples serve to illustrate the invention without limiting its scope.

Temperatures are measured in degrees Celsius. Unless otherwise indicated, the reactions take place at RT.

The R_f values which indicate the ratio of the distance moved by each substance to the distance moved by the eluent front are determined on silica gel thin-layer plates (Merck, Darmstadt, Germany) by thin-layer chromatography using the respective named solvent systems.

Analytical HPLC conditions:

System 1

Linear gradient 20-100% CH_3CN (0.1% TFA) and H_2O (0.1% TFA) in 7 min + 2 min 100% CH_3CN (0.1% TFA); detection at 215 nm, flow rate 1 mL/min at 30°C. Column: Nucleosil 100-3 C18HD (125 x 4mm)

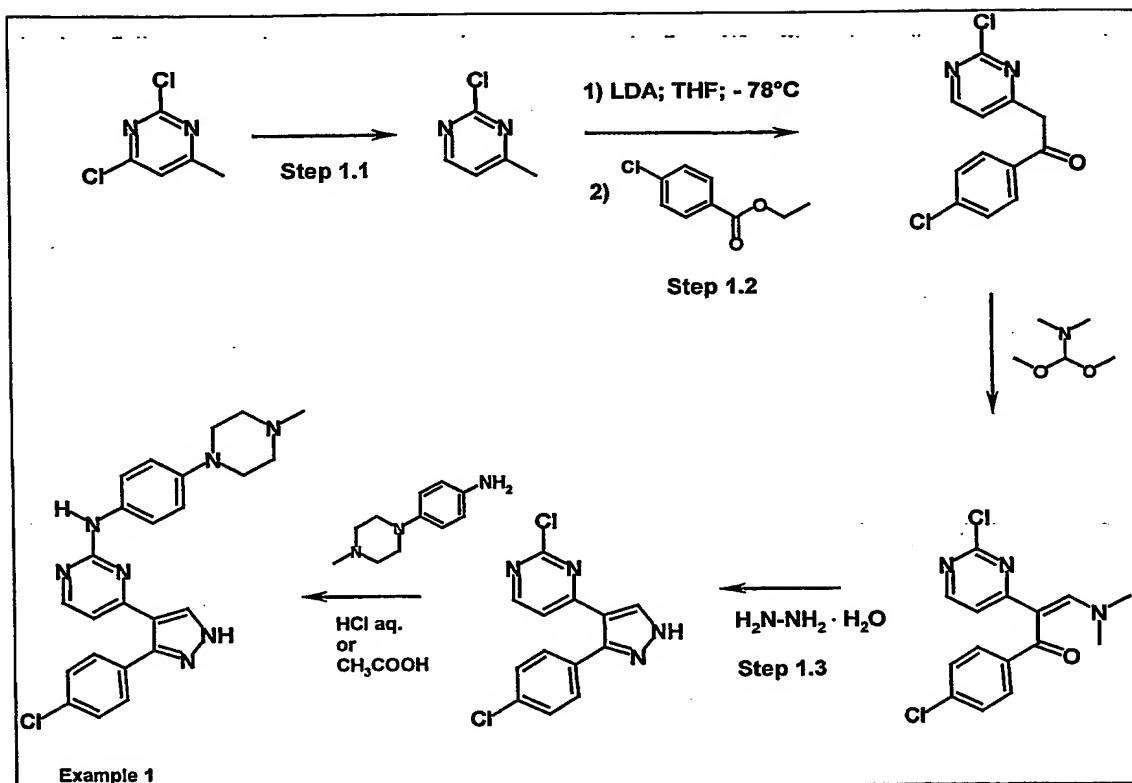
System 2

Linear gradient 2-100% CH_3CN (0.1% TFA) and H_2O (0.1% TFA) in 10 min + 2 min 100% CH_3CN (0.1% TFA); detection at 216 nm, flow rate 2 mL/min at rt. Column: Nucleosil 100-3 C18HD (125 x 4mm)

Abbreviations:

abs.	absolute
approx.	approximately
aq.	aqueous
BOP	(benzotriazol-1-yloxy)-tris(dimethylamino)phosphonium-hexafluoro-phosphate
b.p.	boiling point
DMF	N,N-dimethylformamide
Eq.	equivalent(s)
ES-MS	electron spray-mass spectroscopy
h	hour(s)
HOAc	acetic acid
<i>m</i> -CPBA	3-chloroperoxybenzoic acid
MeOH	methanol
min	minute(s)
m.p.	melting point
rt	room temperature
satd.	saturated
THF	tetrahydrofuran
t _R	retention times

General Synthetic Scheme 1:



Example 1: {4-[3-(4-Chloro-phenyl)-1H-pyrazol-4-yl]pyrimidin-2-yl}-[4-(4-methyl-piperazin-1-yl)-phenyl]-amine

The solution of 150 mg (0.515 mMol) of 2-chloro-4-[3-(4-chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidine in 5 mL of 1,4-dioxane (Merck p.a. 9671) at rt is treated with 99 mg of 4-(4-methyl-piperazin-1-yl)-phenylamine (0.515 mMol, 1 Eq.) and 235 mg (2.06 mMol; 202 μ L; 4 Eq.) of HCl (32%) (Fluka puriss 84416). The yellow suspension is stirred for 24 h at 100°C. After cooling to rt, the supernatant is decanted and discarded. The smearable residue is treated with 10 mL of NaHCO₃ satd. solution and 10 mL of ethyl acetate. The precipitated crude product is filtered off and dried at 60°C under reduced pressure. The combined organic layers are re-extracted with water, dried (Na₂SO₄), and evaporated to obtain additional crude product. The combined crude products are purified by means of column chromatography over silica gel (Si60 (0,040-0,063mm) Merck); eluting with CH₂Cl₂/CH₃OH/NH₄OH (9:1:0.1). Final purification is done by suspending the obtained beige crystals in CH₂Cl₂/CH₃OH (1:1)

to obtain {4-[3-(4-chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(4-methyl-piperazin-1-yl)-phenyl]-amine as pure crystals.

Title compound: m.p.: 282-284°C; ES-MS: 444.3 [M-H]⁺; single peak at t_R = 3.45 min (System 1); R_f = 0.37 (CH₂Cl₂/MeOH/NH₃ 90:10:1).

4-(4-Methyl-piperazin-1-yl)-phenylamine is prepared as follows:

Step A: 50 g (0.247 Mol) of 1-Bromo-4-nitro-benzene and 55 mL (0.495 Mol) of 1-methyl-piperazine are heated for 26 h at 80°C. After cooling, the reaction mixture is taken up in water and extracted with CH₂Cl₂/CH₃OH (8:2). The combined organic layers are dried (Na₂SO₄), filtered and evaporated under reduced pressure. The crude product is recrystallized from ethanol to obtain 1-methyl-4-(4-nitro-phenyl)-piperazine. Title compound: ES-MS: 443.0 [M+H]⁺; R_f = 0.38 (CH₂Cl₂/MeOH 9:1).

Step B: 44.3 g (0.2 Mol) of 1-Methyl-4-(4-nitro-phenyl)-piperazine is dissolved in 1200 mL of CH₃OH and subjected to catalytic hydration at rt using Raney-Ni (10 g) as catalyst. After filtration over Celite, the crude product is purified via solid-distillation (0.16 mbar, 180°C; heat temp 125°C) to obtain 4-(4-methyl-piperazin-1-yl)-phenylamine. Title compound: ES-MS: 192.1 [M+H]⁺; single peak at t_R = 1.08 min (System 1); R_f = 0.33 (CH₂Cl₂/MeOH 4:1).

Step 1.1: 2-Chloro-4-methyl-pyrimidine

200 g (227 Mol) of 2,4-Dichloro-6-methylpyrimidine is suspended in 2 l of water/ethanol (1:1) and heated to 50°C under stirring. Upon dissolution, 331.3 g (5.07 Mol; 4.13 Eq.) of zinc dust (Fluka 96454) is added, followed by 10 crystals of iodine (Merck p.a. 4761.0100). After stirring for 20 h at 50°C, the grey suspension is filtered over HYFLO (Hyflo Super Cel®; Fluka), washed with few ethanol and diluted with 800 mL of water. The mixture is extracted with tert-butylmethyl ether, the organic layer washed with brine, dried (Na₂SO₄), filtered and evaporated under reduced pressure at 40°C to obtain 2-chloro-4-methyl-pyrimidine as beige crystals.

Title compound: m.p.: 44-47°C; single peak at t_R = 2.92 min (System 1); R_f = 0.5 (ethyl acetate).

Step 1.2: 1-(4-Chloro-phenyl)-2-(2-chloro-pyrimidin-4-yl)-ethanone

To the solution of 19 mL (38 mMol) of LDA (Lithiumdiisopropylamide) (~2 M in THF/heptan /ethylbenzene; FLUKA pract 62491) at -74 °C, under a flow of Argon, is added dropwise a

solution of 2-chloro-4-methyl-pyrimidine (4.11 g; 32 mMol) in 20 mL of THF abs. After stirring the beige solution at -74°C for 3 h, 7.01 g (38 mMol) of ethyl-4-chlorobenzoate (LANCASTER 1407) is added within 30 min. The reaction is continued for 21 h, allowing temperature to reach rt. 100 mL of water is added slowly followed by extraction with ethyl acetate. The combined organic layers are washed with water, dried (Na_2SO_4) and filtered. After removal of the solvent under reduced pressure, crude product is obtained. The crude product is suspended in ethyl acetate/hexanes (1:2). After filtering off and drying at 60°C, the orange crystals are recrystallized from methanol to obtain pure, yellowish 1-(4-chlorophenyl)-2-(2-chloro-pyrimidin-4-yl)-ethanone.

Title compound: m.p.: 145-146°C; ES-MS: 267 $[\text{M}+\text{H}]^+$; single peak at $t_R = 6.33$ min (System 1).

Step 1.3: 2-Chloro-4-[3-(4-chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidine

3.67 g (13.74 mMol) of 1-(4-Chloro-phenyl)-2-(2-chloro-pyrimidin-4-yl)-ethanone is suspended in 91.6 mL (50 Eq.) of N,N-dimethylformamid-dimethylacetal (Fluka puriss. 40271) and stirred for 2 h at 100°C. The dark brownish solution is freed from solvent under high vacuum at 30°C. The crude product, (Z)-1-(4-chloro-phenyl)-2-(2-chloro-pyrimidin-4-yl)-3-dimethylamino-propenone, is used for the next step without purification.

To the solution of 5.42 g (14.2 mMol) of (Z)-1-(4-chloro-phenyl)-2-(2-chloro-pyrimidin-4-yl)-3-dimethylamino-propenone in 30 mL of ethanol 0.785 g (0.763 mL; 15.69 mMol, 1.1 Eq.) of hyrazine monohydrate (Fluka 53850) is added at rt. After stirring 45 min at rt, the yellow crystals are filtered off, washed with ethanol and dried at 60°C under vacuum. The crude product is purified by recrystallization from methanol to obtain 2-chloro-4-[3-(4-chlorophenyl)-1H-pyrazol-4-yl]-pyrimidine as slightly yellowish crystals.

Title compound: m.p.: 246-247°C (decomposition temperature); ES-MS: 289 $[\text{M}+\text{H}]^+$; single peak at $t_R = 5.81$ min (System 1); $R_f = 0.2$ ($\text{CH}_2\text{Cl}_2/\text{MeOH}$ 95:5).

Example 2: {4-[3-(4-Chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(2-di methyl-amino-ethoxy)-phenyl]-amine

The title compound is prepared as described in Example 1 using 4-(2-dimethylaminoethoxy)-phenylamine.

4-(2-Dimethylamino-ethoxy)-phenylamine is prepared in 3 steps from *p*-nitrophenol:

Step A: To the solution of 27.83 g (0.2 Mol) of 4-nitro-phenol (Fluka 73560) in 420 mL of acetone is added 55.28 g (0.4 Mol) of potassium carbonate, 143.42 g (1 Mol) of 1-bromo-2-chloro-ethane, 0.55 g (0.0033 Mol) of potassium iodide and 0.28 g (0.00087 Mol) of tetrabutyl-ammonium bromide (Fluka 86860). The resulting suspension is refluxed for 67 h. After removing the solvent under reduced pressure, the residue is taken up into ethyl acetate and washed with water. The combined organic layers are dried (Na_2SO_4), filtered and concentrated under reduced pressure. After trituration of the residue with ligroin, the crystals are filtered off to obtain 1-(2-chloro-ethoxy)-4-nitro-benzene.

Step B: 36 g (0.178 Mol) of 1-(2-Chloro-ethoxy)-4-nitro-benzene is dissolved in 360 mL of ethanol and subjected to catalytic hydration at rt using PtO_2 (1.5 g) as catalyst. The resulting suspension is diluted with CH_2Cl_2 , filtered, and concentrated to approx. 150 mL. After cooling to 0°C the crystals are filtered off, washed and dried at 60°C under vacuum to obtain 4-(2-chloro-ethoxy)-phenylamine. Title compound: m.p.: 87-91°C; ES-MS: 172 $[\text{M}+\text{H}]^+$; single peak at $t_{\text{R}} = 2.73$ min (System 1).

Step C: 11.15 g (0.065 Mol) of 4-(2-Chloro-ethoxy)-phenylamine is suspended in 150 mL (1.18 Mol) of dimethylamine (40% in water; Fluka 38940) and heated under stirring in a steel pressure reactor at 4 bar for 21 h. After cooling the reaction mixture is diluted with 150 mL of 2N NaOH and extracted with ethyl acetate. The combined organic layers are washed with water, dried (Na_2SO_4), filtered and evaporated under reduced pressure to obtain 4-(2-dimethylamino-ethoxy)-phenylamine. Title compound: ES-MS: 181 $[\text{M}+\text{H}]^+$; single peak at $t_{\text{R}} = 1.10$ min (System 1).

Example 3: {4-[3-(4-Chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(2-diethylamino-ethoxy)-phenyl]-amine

The title compound is prepared as described in Example 1 using 4-(2-diethylamino-ethoxy)-phenylamine.

4-(2-Diethylamino-ethoxy)-phenylamine is prepared as follows:

Step A: To the solution of 8.0 g (51 mMol) of 1-chloro-4-nitrobenzene and 6.0 g (6.8 mL; 51 mMol) of 2-diethylamino-ethanol in 50 mL of DMF at 0°C is added 2.7 g of NaH portionwise over 2 h. After stirring another hour at rt, the reaction mixture is poured onto 300 mL of water and stirred. After filtering off some crystalline product, the aqueous layers are extracted with

ethyl acetate. The combined organic layers are dried and evaporated to obtain diethyl-[2-(4-nitro-phenoxy)-ethyl]-amine. Title compound: ES-MS: 239 [M+H]⁺; single peak at t_R = 5.1 min (System 2).

Step B: 10 g (42 mMol) of Diethyl-[2-(4-nitro-phenoxy)-ethyl]-amine is dissolved in 100 mL of ethanol and subjected to catalytic hydration at rt using 894 mg of Pd/C. After filtration through HYFLO and removal of the solvent under reduced pressure, the crude product is purified by bulb-to-bulb distillation (140°C) to obtain 4-(2-diethylamino-ethoxy)-phenylamine. Title compound: ES-MS: 209 [M+H]⁺; single peak at t_R = 2.3 min (System 2).

Example 4: {4-[3-(4-Chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(2-morpholin-4-yl-ethoxy)-phenyl]-amine

The title compound is prepared as described in Example 1 using 4-(2-morpholin-4-yl-ethoxy)-phenylamine.

4-(2-Morpholin-4-yl-ethoxy)-phenylamine is prepared as follows:

Step A: To the solution of 8.0 g (51 mMol) of 1-chloro-4-nitrobenzene and 6.7 g (6.3 mL; 51 mMol) of N-hydroxyethyl-morpholine in 50 mL of DMF at 0°C is added 2.7 g of NaH portionwise over 2.5 h. After stirring another hour at rt, the reaction mixture is poured onto 200 mL of water and stirred. The precipitated crystals are filtered and dried at 60°C under vacuum to obtain 4-[2-(4-nitro-phenoxy)-ethyl]-morpholine. Title compound: ES-MS: 253 [M+H]⁺; single peak at t_R = 4.8 min (System 2).

Step B: 8.1 g (32 mMol) of 4-[2-(4-Nitro-phenoxy)-ethyl]-morpholine is dissolved in 100 mL of ethanol and subjected to catalytic hydration at rt using 681 mg of Pd/C. After filtration through HYFLO and removal of the solvent under reduced pressure, the crude product is purified by bulb-to-bulb distillation (0.13 mbar; 200°C) to obtain 4-(2-morpholin-4-yl-ethoxy)-phenylamine. Title compound: ES-MS: 223 [M+H]⁺; single peak at t_R = 3 min (System 2).

Example 5: {4-[3-(3-Chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(2-dimethylamino-ethoxy)-phenyl]-amine

The title compound is prepared as described in Example 1 using 3-chloro-benzoic acid ethyl ester and 4-(2-dimethylamino-ethoxy)-phenylamine (see Example 2).

3-Chloro-benzoic acid ethyl ester is prepared from 3-chloro-benzoic acid using a standard protocol with ethanol/H₂SO₄ according to L.-F. Tietze & T. Eicher, *Reactions and Syntheses in the Organic Chemistry Laboratory*; University Science Books, Mill Valley, CA, 1989). Title compound: single peak at t_R= 7.29 min (System 1).

Example 6: {4-[3-(3-Chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(2-morpholin-4-yl-ethoxy)-phenyl]-amine

The title compound is prepared as described in Example 1 using 3-chloro-benzoic acid ethyl ester (see Example 5) and 4-(2-morpholin-4-yl-ethoxy)-phenylamine (see Example 4).

Example 7: {4-[3-(3-Chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(2-diethylamino-ethoxy)-phenyl]-amine

The title compound is prepared as described in Example 1 using 3-chloro-benzoic acid ethyl ester (see Example 5) and 4-(2-diethylamino-ethoxy)-phenylamine (see Example 3).

Example 8: {4-[3-(3-Chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(4-methyl-piperazin-1-yl)-phenyl]-amine

The title compound is prepared as described in Example 1 using 3-chloro-benzoic acid ethyl ester (see Example 5).

Example 9: {4-[3-(4-Methoxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(4-methyl-piperazin-1-yl)-phenyl]-amine

The title compound is prepared as described in Example 1 using 4-methoxy-benzoic acid ethyl ester (Aldrich W24,200-4).

Example 10: {4-[3-(4-Methoxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(2-morpholin-4-yl-ethoxy)-phenyl]-amine

The title compound is prepared as described in Example 1 using 4-methoxy-benzoic acid ethyl ester (Aldrich W24,200-4) and 4-(2-morpholin-4-yl-ethoxy)-phenylamine (see Example 4).

Example 11: {4-[3-(4-Ethyl-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(4-methyl-piperazin-1-yl)-phenyl]-amine

The title compound is prepared as described in Example 1 using 4-ethyl-benzoic acid ethyl ester.

4-Ethyl-benzoic acid ethyl ester is prepared from 4-ethyl-benzoic acid using a standard protocol with ethanol/H₂SO₄ according to L.-F. Tietze & T. Eicher, *Reactions and Syntheses in the Organic Chemistry Laboratory*; University Science Books, Mill Valley, CA, 1989). Title compound: b.p. 132°C (12 mm Hg); single peak at *t*_R = 7.51 min (System 1).

Example 12: [4-(2-Diethylamino-ethoxy)-phenyl]-{4-[3-(4-ethyl-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-amine

The title compound is prepared as described in Example 1 using 4-ethyl-benzoic acid ethyl ester (see Example 11) and 4-(2-diethylamino-ethoxy)-phenylamine (see Example 3).

Example 13: [4-(2-Diethylamino-ethoxy)-phenyl]-{4-[3-(4-methoxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-amine

The title compound is prepared as described in Example 1 using 4-methoxy-benzoic acid ethyl ester (Aldrich W24,200-4) and 4-(2-diethylamino-ethoxy)-phenylamine (see Example 3).

Example 14: [4-(2-Diethylamino-ethoxy)-phenyl]-{4-[3-(3-methoxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-amine

The title compound is prepared as described in Example 1 using 3-methoxy-benzoic acid ethyl ester and 4-(2-diethylamino-ethoxy)-phenylamine (see Example 3).

3-Methoxy-benzoic acid ethyl ester is prepared from 3-methoxy-benzoic acid using a standard protocol with ethanol/H₂SO₄ according to L.-F. Tietze & T. Eicher, *Reactions and Syntheses in the Organic Chemistry Laboratory*; University Science Books, Mill Valley, CA, 1989). Title compound: b.p. 151°C (25 mm Hg); single peak at *t*_R = 6.51 min (System 1).

Example 15: {4-[3-(3-Methoxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(2-morpholin-4-yl-ethoxy)-phenyl]-amine

The title compound is prepared as described in Example 1 using 3-methoxy-benzoic acid ethyl ester (see Example 14) and 4-(2-morpholin-4-yl-ethoxy)-phenylamine (see Example 4).

Example 16: {4-[3-(3-Methoxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(4-methyl-piperazin-1-yl)-phenyl]-amine

The title compound is prepared as described in Example 1 using 3-methoxy-benzoic acid ethyl ester (see Example 14).

Example 17: [4-(2-Dimethylamino-ethoxy)-phenyl]-[4-[3-(4-ethyl-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl]-amine

The title compound is prepared as described in Example 1 using 4-ethyl-benzoic acid ethyl ester (see Example 11) and 4-(2-dimethylamino-ethoxy)-phenylamine (see Example 2).

Example 18: [4-(4-Methyl-piperazin-1-yl)-phenyl]-[4-(3-m-tolyl-1H-pyrazol-4-yl)-pyrimidin-2-yl]-amine

The title compound is prepared as described in Example 1 using ethyl-3-methylbenzoate (Aldrich 25,305-7).

Example 19: [4-(2-Diethylamino-ethoxy)-phenyl]-[4-(3-m-tolyl-1H-pyrazol-4-yl)-pyrimidin-2-yl]-amine

The title compound is prepared as described in Example 1 using ethyl-3-methylbenzoate (Aldrich 25,305-7) and 4-(2-diethylamino-ethoxy)-phenylamine (see Example 3).

Example 20: [4-(2-Dimethylamino-ethoxy)-phenyl]-[4-(3-m-tolyl-1H-pyrazol-4-yl)-pyrimidin-2-yl]-amine

The title compound is prepared as described in Example 1 using ethyl-3-methylbenzoate (Aldrich 25,305-7) and 4-(2-dimethylamino-ethoxy)-phenylamine (see Example 2).

Example 21: {4-[3-(3,4-Dichloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(2-dimethylamino-ethoxy)-phenyl]-amine

The title compound is prepared as described in Example 1 using ethyl-3,4-dichlorobenzoate and 4-(2-dimethylamino-ethoxy)-phenylamine (see Example 2).

Ethyl-3,4-dichlorobenzoate is prepared from 3,4-dichloro-benzoic acid using a standard protocol with ethanol/H₂SO₄ according to L.-F. Tietze & T. Eicher, Reactions and Syntheses in the Organic Chemistry Laboratory; University Science Books, Mill Valley, CA, 1989). Title compound: m.p.: 40-41°C; single peak at t_R = 7.89 min (System 1).

Example 22: {4-[3-(3,4-Dichloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(4-methyl-piperazin-1-yl)-phenyl]-amine

The title compound is prepared as described in Example 1 using ethyl-3,4-dichlorobenzoate (see Example 21).

Example 23: {4-[3-(4-Benzyl-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(2-dimethylamino-ethoxy)-phenyl]-amine

The title compound is prepared as described in Example 1 using ethyl-4-(benzyloxy)benzoate (MAYBRIDGE 03-1741) and 4-(2-dimethylamino-ethoxy)-phenylamine (see Example 2).

Example 24: 4-(4-{2-[4-(4-Methyl-piperazin-1-yl)-phenylamino]-pyrimidin-4-yl}-1H-pyrazol-3-yl)-phenol

The title compound is prepared as described in Example 1 using ethyl-4-(benzyloxy)benzoate (MAYBRIDGE 03-1741). The title compound is isolated as a side product due to the loss of the benzyl group.

Example 25: [4-(4-Methyl-piperazin-1-yl)-phenyl]-[4-(3-p-tolyl-1H-pyrazol-4-yl)-pyrimidin-2-yl]-amine

The title compound is prepared as described in Example 1 using ethyl-p-toluate (Fluka 89909).

Example 26: [4-(2-Dimethylamino-ethoxy)-phenyl]-[4-(3-p-tolyl-1H-pyrazol-4-yl)-pyrimidin-2-yl]-amine

The title compound is prepared as described in Example 1 using ethyl-4-methylbenzoate (see Example 25) and 4-(2-dimethylamino-ethoxy)-phenylamine (see Example 2).

Example 27: {4-[3-(4-Chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[3-(1-methyl-piperidin-4-ylmethoxy)-phenyl]-amine

The title compound is prepared as described in Example 1 using [3-(1-methyl-piperidin-4-ylmethoxy)-phenyl]-carbamic acid tert-butyl ester.

[3-(1-Methyl-piperidin-4-ylmethoxy)-phenyl]-carbamic acid tert-butyl ester is prepared as follows:

To the solution of 3.04 g (14.51 mMol; 1.25 Eq.) of 3-N-BOC-Aminophenol and 3.81 g (14.51 mMol; 1.25 Eq.) of triphenylphosphine (Aldrich T8,440-9) under Argon in 30 mL of THF at 10°C is added dropwise a solution of 2.26 mL (14.51 mMol) of diethyl-azodicarboxylate (95%; Fluka 11624) in 6 mL of THF abs. After stirring for 10 min under ice cooling, a solution of 1.5 g (11.61 mMol; 1 Eq.) of 1-methyl-4-piperidinemethanol (Chem Pacific; 33077*) in 6 mL of THF abs. is added and kept for 20 min at 10°C. After 15 h at rt, the solvent is removed under reduced pressure and the reaction mixture is purified by column chromatography over silica gel [Si60 (0.040-0.063mm) Merck], eluting with methylenchlorid/methanol/NH₃ (25% aqua) 70:10:0.8 to obtain [3-(1-methyl-piperidin-4-ylmethoxy)-phenyl]-carbamic acid tert-butyl ester. Title compound: ES-MS: 321.1 [M+H]⁺; single peak at t_R = 4.63 min (System 1).

* 1-Methyl-4-piperidinemethanol can alternatively be prepared from piperidin-4-yl-methanol and formaldehyde (36% in water) under reductive conditions (H₂/RaNi in CH₃OH) at rt.

Example 28: {4-[3-(4-Chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(1-methyl-piperidin-4-ylmethoxy)-phenyl]-amine

The title compound is prepared as described in Example 1 using [4-(1-methyl-piperidin-4-ylmethoxy)-phenyl]-carbamic acid tert-butyl ester.

[4-(1-Methyl-piperidin-4-ylmethoxy)-phenyl]-carbamic acid tert-butyl ester is prepared as described for [3-(1-methyl-piperidin-4-ylmethoxy)-phenyl]-carbamic acid tert-butyl ester (see Example 27) using 4-N-BOC-Aminophenol (AstaTech; B56686). Title compound: ES-MS: 321.1 [M+H]⁺; single peak at t_R = 4.55 min (System 1).

Example 29: {4-[3-(4-Chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[3-(4-methyl-piperazin-1-yl)-phenyl]-amine

The title compound is prepared as described in Example 1 using 3-(4-methyl-piperazin-1-yl)-phenylamine.

3-(4-Methyl-piperazin-1-yl)-phenylamine is prepared as described for 4-(4-methyl-piperazin-1-yl)-phenylamine (see Example 1) using 3-bromo-phenylamine.

Title compound: ES-MS: 192.0 [M+H]⁺; single peak at t_R = 1.14 min (System 1).

Example 30: 4-[4-[3-(4-Chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-ylamino]-benzoic acid

The title compound is prepared as described in Example 1 using 4-aminobenzoic acid (Fluka 06930).

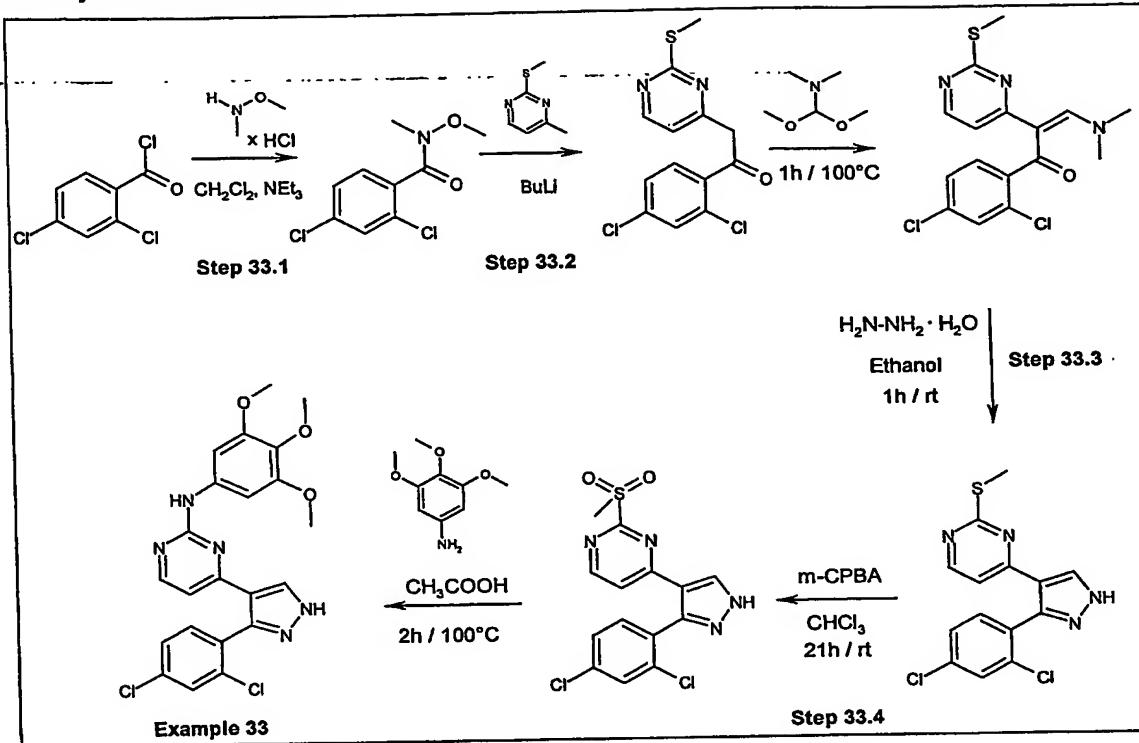
Example 31: (4-{4-[3-(4-Chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-ylamino}-phenyl)-(4-methyl-piperazin-1-yl)-methanone

To the suspension of 600 mg (1.53 mMol; 1 Eq.) of 4-{4-[3-(4-chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-ylamino}-benzoic acid (Example 30) in 5 mL of DMF at rt is added 186 mg (202 μ L; 1.83 mMol; 1.2 Eq.) of N-methylmorpholine (Fluka 67869); 677 mg (1.53 mMol; 1 Eq.) of BOP (Fluka 12802) and 153 mg (170 μ L; 1.53 mMol; 1 Eq.) of N-methylpiperazine (Fluka 68810). After 2 h at rt, the yellow solution is treated with 50 mL of NaHCO_3 satd. solution and extracted with ethyl acetate. The combined organic layers are washed with water, dried (Na_2SO_4), filtered and concentrated under reduced pressure to approx. 10 mL. The precipitated crystals are collected, washed with a little ethyl acetate and dried at 60°C under vacuum to obtain (4-{4-[3-(4-chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-ylamino}-phenyl)-(4-methyl-piperazin-1-yl)-methanone.

Example 32: {4-[3-(4-Chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(4-methyl-piperazin-1-ylmethyl)-phenyl]-amine

To the suspension of 96 mg (2.53 mMol; 4 Eq.) of lithium aluminum hydride (Fluka 62420) in 20 mL of THF abs. under Argon at rt is added slowly within 10 min 300 mg (0.633 mMol) of (4-{4-[3-(4-chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-ylamino}-phenyl)-(4-methyl-piperazin-1-yl)-methanone (see Example 31) as a solid; under vigorous foaming. After stirring for 2.5 h at rt, the suspension is cooled to 0°C and treated dropwise with 20 mL of water. After diluting further with 50 mL of water, the suspension is extracted with ethyl acetate. The combined organic layers are washed with water, dried (Na_2SO_4), filtered and concentrated under reduced pressure. The crude product is purified over silica gel, eluting with $\text{CH}_2\text{Cl}_2/\text{CH}_3\text{OH}$ (4:1) followed by trituration in diethyl ether to obtain {4-[3-(4-chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(4-methyl-piperazin-1-ylmethyl)-phenyl]-amine.

General Synthetic Scheme 2:



Example 33: {4-[3-(2,4-Dichloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-(3,4,5-trimethoxy-phenyl)-amine

{4-[3-(2,4-Dichloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-(3,4,5-trimethoxy-phenyl)-amine is synthesized from 4-[3-(2,4-dichloro-phenyl)-1H-pyrazol-4-yl]-2-methanesulfonyl-pyrimidine in analogy to a literature procedure (Klutchko et al., Journal of Medicinal Chemistry, 1998, Vol. 41, No. 17, 3276-3292) using 3,4,5-trimethoxyaniline (Fluka 92129) and glacial HOAc.

Step 33.1: 2,4-Dichloro-N-methoxy-N-methyl-benzamide

The title compound is prepared in analogy to a standard protocol using 2,4-dichloro-benzoyl chloride (Aldrich 11,193-7) and N-O-dimethylhydroxylamine HCl (Fluka 40706) in CH_2Cl_2 (Nahm, Steven; Weinreb, Steven M.; *Tetrahedron Lett.*; 1981; 22 (39); 3815-3818).

Step 33.2: 1-(2,4-Dichloro-phenyl)-2-(2-methylsulfanyl-pyrimidin-4-yl)-ethanone

To the solution of 85 mL of diisopropylamine (Fluka 38300) and 2.5 L of THF at -75°C under Argon is added 370 mL (0.55 Mol) of butyl lithium (BuLi; 1.6N in hexanes; Fluka 20160) within 1 h. To this is added 71 g (0.5 Mol) of 4-methyl-2-methylsulfanyl-pyrimidine in 250 mL of CH_2Cl_2 at -75°C within 30 min. After this, a solution of 118 g (0.5 Mol) of 2,4-dichloro-N-

methoxy-N-methyl-benzamide in 250 mL of CH_2Cl_2 at -75°C within 30 min followed by letting the mixture warm up to rt. After completion, the reaction mixture is poured onto 7 L of NH_4Cl satd. and extracted with ethyl acetate. The combined organic layers are washed with brine, dried and evaporated. The crude product is suspended in hexanes. After filtering off and drying 1-(2,4-dichloro-phenyl)-2-(2-methylsulfanyl-pyrimidin-4-yl)-ethanone is obtained as yellow crystals. Title compound: m.p.: 105-107°C; ES-MS: 313.0 / 314.9 $[\text{M}+\text{H}]^+$.

4-Methyl-2-methylsulfanyl-pyrimidine is prepared from 4-methyl-pyrimidine-2-thiol in analogy to a standard protocol using methyliodide (Strekowski, L.; Wydra, R. L.; Janda, L.; Harden, D. B. *J. Org. Chem.* (1991), 56(19), 5610-14). Title compound: single peak at 3.94 min (System 1); $R_f = 0.32$ (ethyl acetate / hexane 1:2).

Step 33.3: 4-[3-(2,4-Dichloro-phenyl)-1H-pyrazol-4-yl]-2-methylsulfanyl-pyrimidine

The title compound is prepared from 1-(2,4-dichloro-phenyl)-2-(2-methylsulfanyl-pyrimidin-4-yl)-ethanone as described in Example 1, Step 1.3, using N,N-dimethylformamid-dimethylacetal and hydrazine monohydrate. Title compound: m.p.: 218-219°C; single peak at 6.07 min (System 1).

Step 33.4: 4-[3-(2,4-Dichloro-phenyl)-1H-pyrazol-4-yl]-2-methanesulfonyl-pyrimidine

The title compound is prepared from 4-[3-(2,4-dichloro-phenyl)-1H-pyrazol-4-yl]-2-methylsulfanyl-pyrimidine in analogy to a literature procedure (Klutchko et al., *Journal of Medicinal Chemistry*, 1998, Vol. 41, No. 17, 3276-3292) using *m*-CPBA (Fluka 25800) in CHCl_3 . Title compound: ES-MS: 368.9 $[\text{M}+\text{H}]^+$; single peak at $t_R = 4.98$ min (System 1); $R_f = 0.38$ (ethyl acetate).

Example 34: {4-[3-(2,4-Dichloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(4-methyl-piperazin-1-yl)-phenyl]-amine

The title compound is prepared as described in Example 33 using 4-(4-methyl-piperazin-1-yl)-phenylamine (see Example 1).

Example 35: N-{4-[3-(4-Chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-N',N'-dimethyl-benzene-1,3-diamine

The title compound is prepared as described in Example 1 using 3-amino-N,N-dimethylaniline dihydrochloride (Fluka 07765).

Example 36: {4-[3-(4-Ethoxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(4-methyl-piperazin-1-yl)-phenyl]-amine

The title compound is prepared as described in Example 33 using ethyl-4-ethoxybenzoate 97% (ACROS 346590250) and 4-(4-methyl-piperazin-1-yl)-phenylamine (see Example 1).

Example 37: {4-[3-(4-Ethoxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-(3,4,5-trimethoxy-phenyl)-amine

The title compound is prepared as described in Example 33 using 4-ethoxy-benzoic acid ethyl ester (ACROS 346590250) and 3,4,5-trimethoxyaniline (Fluka 92129).

Physicochemical data:

Example	melting point from / to	HPLC [min] / System	mass spectra; ESI; m/z =
1	282	284	3.45 / 1 444.3 [M-H] ⁻
2	236	239	3.48 / 1 435 [M+H] ⁺
3	191	192	3.74 / 1 463 [M+H] ⁺
4	244	247	3.53 / 1 477 [M+H] ⁺
5	182	183	3.46 / 1 435 [M+H] ⁺
6	192	193	3.54 / 1 477 [M+H] ⁺
7	179	181	3.74 / 1 463 [M+H] ⁺
8	297	299	3.43 / 1 446 [M+H] ⁺
9	273	277	3.15 / 1 442.0 [M+H] ⁺
10	232	234	3.23 / 1 473.0 [M+H] ⁺
11	282	283	3.75 / 1 457.2 [M+H] ⁺
12	206	207	4.01 / 1 440.2 [M+H] ⁺
13		3.44 / 1	459.0 [M+H] ⁺
14		3.48 / 1	459.3 [M+H] ⁺
15		3.28 / 1	473.2 [M+H] ⁺
16		3.17 / 1	442.2 [M+H] ⁺
17	242	244	3.77 / 1 429.1 [M+H] ⁺
18	280	282	3.38 / 1 426.1 [M+H] ⁺
19	181	182	3.68 / 1 440.2 [M+H] ⁺

20	206	207	3.41 / 1	415.1 [M+H] ⁺
21	188	191	3.73 / 1	469.0 [M+H] ⁺
22	300	307	3.70 / 1	479.9 [M+H] ⁺
23	213	216	3.63 / 1	507.0 [M+H] ⁺
24	305	309	2.40 / 1	428.1 [M+H] ⁺
25	289	294	3.43 / 1	426.0 [M+H] ⁺
26	251	259	3.45 / 1	415.1 [M+H] ⁺
27	225	228	4.15 / 1	475.0 [M+H] ⁺
28	270	274	3.73 / 1	475.0 [M+H] ⁺
29	274	277	3.82 / 1	446.0 [M+H] ⁺
30	342	349	4.53 / 1	391.9 [M+H] ⁺
31	173	175	3.63 / 1	473.9 [M+H] ⁺
32	240	242	3.41 / 1	460.0 [M+H] ⁺
33	125	127	4.96 / 1	472.0 [M+H] ⁺
34	263	265	3.52 / 1	477.9 [M+H] ⁺
35			3.90 / 1	391.2 [M+H] ⁺
36			3.37 / 1	456.2 [M+H] ⁺
37	293	297	4.71 / 1	

Examples 38 – 49:

The following Examples are synthesized using an analogous procedure as described in Example 33:

Example Number	Name
38	{4-[3-(4-Chloro-3-methoxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-(3,4,5-trimethoxy-phenyl)-amine
39	3-{4-[3-(4-Chloro-3-hydroxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-ylamino}-benzenesulfonamide
40	3-{4-[3-(4-Chloro-3-methoxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-ylamino}-benzenesulfonamide

41	3-{4-[3-(4-Chloro-3-methoxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-ylamino}-benzoic acid ethyl ester
42	3-{4-[3-(4-Chloro-3-hydroxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-ylamino}-benzoic acid ethyl ester
43	1-(3-{4-[3-(4-Chloro-3-methoxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-ylamino}-phenyl)-ethanone
44	Benzothiazol-6-yl-{4-[3-(4-chloro-3-methoxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-amine
45	{4-[3-(4-Chloro-3-methoxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-(1-oxo-benzothiazol-6-yl)-amine
46	[4-(3-Benzo[1,3]dioxol-5-yl-1H-pyrazol-4-yl)-pyrimidin-2-yl]-(3,4,5-trimethoxy-phenyl)-amine
47	3-[4-(3-Benzo[1,3]dioxol-5-yl-1H-pyrazol-4-yl)-pyrimidin-2-ylamino]-benzoic acid tert-butyl ester
48	3-[4-(3-Benzo[1,3]dioxol-5-yl-1H-pyrazol-4-yl)-pyrimidin-2-ylamino]-benzenesulfonamide
49	[4-(3-Benzo[1,3]dioxol-5-yl-1H-pyrazol-4-yl)-pyrimidin-2-yl]-(3-methylsulfanyl-phenyl)-amine

Example 50: Dry-filled capsules

5000 capsules, each comprising as active ingredient 0.25 g of one of the compounds of formula I mentioned in the preceding Examples, are prepared as follows:

Composition

active ingredient	1250 g
talcum	180 g

wheat starch	120 g
magnesium stearate	80 g
lactose	20 g

Preparation process: The mentioned substances are pulverised and forced through a sieve of 0.6 mm mesh size. 0.33 g portions of the mixture are introduced into gelatin capsules using a capsule-filling machine.

Example 51: Soft capsules

5000 soft gelatin capsules, each comprising as active ingredient 0.05 g of one of the compounds of formula I mentioned in the preceding Examples, are prepared as follows:

Composition

active ingredient	250 g
PEG 400	1 litre
Tween 80	1 litre

Preparation process: The active ingredient is pulverised and suspended in PEG 400 (polyethylene glycol having an M_r of from approx. 380 to approx. 420, Fluka, Switzerland) and Tween[®]80 (polyoxyethylene sorbitan monolaurate, Atlas Chem. Ind. Inc., USA, supplied by Fluka, Switzerland) and ground in a wet pulveriser to a particle size of approx. from 1 to 3 μm . 0.43 g portions of the mixture are then introduced into soft gelatin capsules using a capsule-filling machine.

Example 52: Inhibition of the tyrosine kinase activity of EGF-R (HER-1), ErbB-2 (HER-2) and VEGF receptor (KDR)

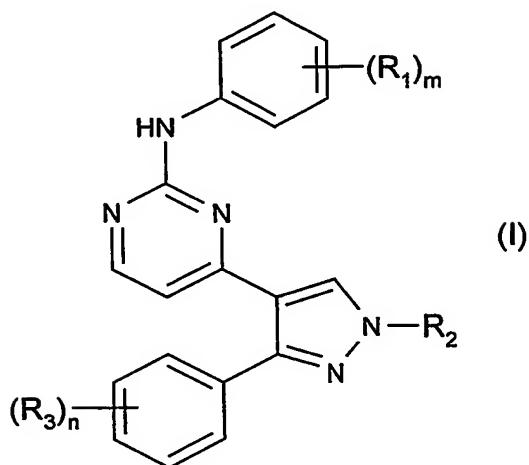
The inhibition tests are carried out as described above. The IC_{50} values for some of the compounds of formula I are given below:

Compound from Example No.	HER-1 IC_{50} [μM]	HER-2 IC_{50} [μM]	KDR IC_{50} [μM]
1	0.018	0.023	0.01
2	0.022	0.019	0.021

18	0.021	0.013	0.004
20	0.027	0.015	0.006
21	0.029	0.016	0.015
46	0.028	0.006	0.003

What is claimed is:

1. A compound of formula I



wherein

m and n are independently of one another an integer from 1 to 5;

R₁ is unsubstituted, mono- or di-substituted amino-sulfonyl; unsubstituted, mono- or di-substituted amino; a heterocyclic radical; lower alkyl substituted by a heterocyclic radical; a radical R₄-lower alkyl-X-, wherein R₄ is hydrogen, unsubstituted, mono- or di-substituted amino, or a heterocyclic radical, and X is -S- or -O-; or a radical R₅-C(=O)-, wherein R₅ is hydrogen, unsubstituted or substituted lower alkyl, free or etherified hydroxy, unsubstituted, mono- or di-substituted amino, or a heterocyclic radical; wherein the R₁ substituents are selected independently of one another if m>1;

or two vicinal R₁ substituents together with the phenyl carbon atoms to which they are attached form a heterocyclic ring;

R₂ is hydrogen, unsubstituted or substituted lower alkyl or a heterocyclic radical; and

R₃ is halogen; unsubstituted or substituted lower alkyl; or free, etherified or esterified hydroxy; wherein the R₃ substituents are selected independently of one another if n>1; or two vicinal R₃ substituents together with the phenyl carbon atoms to which they are attached form a heterocyclic ring;

with the proviso that R₁ is not methoxy if m and n are both 1, R₂ is hydrogen and R₃ is fluoro; or a salt thereof.

2. A compound of formula I according to claim 1, wherein
m and n are independently of one another an integer from 1 to 5;
R₁ is unsubstituted, mono- or di-substituted amino-sulfonyl; unsubstituted, mono- or di-
substituted amino; a heterocyclic radical; lower alkyl substituted by a heterocyclic radical; a
radical R₄-lower alkyl-X-, wherein R₄ is hydrogen, unsubstituted, mono- or di-substituted
amino, or a heterocyclic radical, and X is -S- or -O-; or a radical R₅-C(=O)-, wherein R₅ is
hydrogen, unsubstituted or substituted lower alkyl, free or etherified hydroxy, unsubstituted,
mono- or di-substituted amino, or a heterocyclic radical; wherein the R₁ substituents are
selected independently of one another if m>1;
or two vicinal R₁ substituents together with the phenyl carbon atoms to which they are
attached form a heterocyclic ring;
R₂ is hydrogen; and
R₃ is halogen; unsubstituted or substituted lower alkyl; or free, etherified or esterified
hydroxy; wherein the R₃ substituents are selected independently of one another if n>1;
or two vicinal R₃ substituents together with the phenyl carbon atoms to which they are
attached form a heterocyclic ring;
with the proviso that R₁ is not methoxy if m and n are both 1, R₂ is hydrogen and R₃ is fluoro;
or a salt thereof.

3. A compound of formula I according to claim 1, wherein
m and n are independently of one another an integer from 1 to 3;
R₁ is amino-sulfonyl; mono- or di-substituted amino; a heterocyclic radical; lower alkyl
substituted by a heterocyclic radical; a radical R₄-lower alkyl-X-, wherein R₄ is hydrogen,
mono- or di-substituted amino, or a heterocyclic radical, and X is -S- or -O-; or a radical R₅-
C(=O)-, wherein R₅ is lower alkyl, free or etherified hydroxy, or a heterocyclic radical;
wherein the R₁ substituents are selected independently of one another if m>1;
or two vicinal R₁ substituents together with the phenyl carbon atoms to which they are
attached form a heterocyclic ring;
R₂ is hydrogen; and
R₃ is halogen; lower alkyl; or free or etherified hydroxy; wherein the R₃ substituents are
selected independently of one another if n>1;
or two vicinal R₃ substituents together with the phenyl carbon atoms to which they are
attached form a heterocyclic ring;
with the proviso that R₁ is not methoxy if m and n are both 1, R₂ is hydrogen and R₃ is fluoro;

or a salt thereof.

4. A compound of formula I according to claim 1, wherein

m is an integer from 1 to 3 and n is 1 or 2;

R_1 is amino-sulfonyl; N,N-di-lower alkylamino; lower alkyl-piperazinyl; lower alkyl substituted by lower alkyl-piperazinyl; a radical R_4 -lower alkyl-X-, wherein R_4 is hydrogen, N,N-di-lower alkylamino, morpholinyl or lower alkyl-piperidyl, and X is -S- or -O-; or a radical R_5 -C(=O)-, wherein R_5 is lower alkyl, hydroxy, lower alkoxy or lower alkyl-piperazinyl; wherein the R_1 substituents are selected independently of one another if $m > 1$;

or two vicinal R_1 substituents together with the phenyl carbon atoms to which they are attached form a thiazol or 1-oxo-thiazol ring;

R_2 is hydrogen; and

R_3 is halogen, lower alkyl, hydroxy, lower alkoxy or phenyl-lower alkoxy; wherein the R_3 substituents are selected independently of one another if n is 2;

or two R_3 radicals together form a dioxol ring;

with the proviso that R_1 is not methoxy if m and n are both 1, R_2 is hydrogen and R_3 is fluoro; or a salt thereof.

5. A compound of formula I according to any one of claims 1, wherein

m and n are independently of one another an integer from 1 to 5;

R_1 is unsubstituted, mono- or di-substituted amino-sulfonyl; unsubstituted, mono- or di-substituted amino; a heterocyclic radical; lower alkyl substituted by a heterocyclic radical; or a radical R_5 -C(=O)-, wherein R_5 is hydrogen, unsubstituted or substituted lower alkyl, free or etherified hydroxy, unsubstituted, mono- or di-substituted amino, or a heterocyclic radical; wherein the R_1 substituents are selected independently of one another if $m > 1$;

or two vicinal R_1 substituents together with the phenyl carbon atoms to which they are attached form a heterocyclic ring;

R_2 is hydrogen, unsubstituted or substituted lower alkyl or a heterocyclic radical; and

R_3 is halogen; unsubstituted or substituted lower alkyl; or free, etherified or esterified hydroxy; wherein the R_3 substituents are selected independently of one another if $n > 1$; or two vicinal R_3 substituents together with the phenyl carbon atoms to which they are attached form a heterocyclic ring;

or a salt thereof.

6. A compound of formula I according to claim 1, selected from the group consisting of
{4-[3-(4-chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(4-methyl-
piperazin-1-yl)-phenyl]-amine;
{4-[3-(4-chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(2-di methyl-amino-ethoxy)-
phenyl]-amine;
{4-[3-(4-chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(2-diethylamino-ethoxy)-phenyl]-
amine;
{4-[3-(4-chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(2-morpholin-4-yl-ethoxy)-phenyl]-
amine;
{4-[3-(3-chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(2-dimethylamino-ethoxy)-phenyl]-
amine;
{4-[3-(3-chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(2-morpholin-4-yl-ethoxy)-phenyl]-
amine;
{4-[3-(3-chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(2-diethylamino-ethoxy)-phenyl]-
amine;
{4-[3-(3-chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(4-methyl-piperazin-1-yl)-phenyl]-
amine;
{4-[3-(4-methoxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(4-methyl-piperazin-1-yl)-
phenyl]-amine;
{4-[3-(4-methoxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(2-morpholin-4-yl-ethoxy)-
phenyl]-amine;
{4-[3-(4-ethyl-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(4-methyl-piperazin-1-yl)-phenyl]-
amine;
{4-(2-diethylamino-ethoxy)-phenyl}-[4-[3-(4-ethyl-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl]-
amine;
{4-(2-diethylamino-ethoxy)-phenyl}-[4-[3-(4-methoxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl]-
amine;
{4-(2-diethylamino-ethoxy)-phenyl}-[4-[3-(3-methoxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl]-
amine;
{4-[3-(3-methoxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(2-morpholin-4-yl-ethoxy)-
phenyl]-amine;
{4-[3-(3-methoxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(4-methyl-piperazin-1-yl)-
phenyl]-amine;

[4-(2-dimethylamino-ethoxy)-phenyl]-{4-[3-(4-ethyl-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-amine;
[4-(4-methyl-piperazin-1-yl)-phenyl]-[4-(3-m-tolyl-1H-pyrazol-4-yl)-pyrimidin-2-yl]-amine;
[4-(2-diethylamino-ethoxy)-phenyl]-[4-(3-m-tolyl-1H-pyrazol-4-yl)-pyrimidin-2-yl]-amine;
[4-(2-dimethylamino-ethoxy)-phenyl]-[4-(3-m-tolyl-1H-pyrazol-4-yl)-pyrimidin-2-yl]-amine;
{4-[3-(3,4-dichloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(2-dimethylamino-ethoxy)-phenyl]-amine;
{4-[3-(3,4-dichloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(4-methyl-piperazin-1-yl)-phenyl]-amine;
{4-[3-(4-benzyloxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(2-dimethylamino-ethoxy)-phenyl]-amine;
4-(4-{2-[4-(4-methyl-piperazin-1-yl)-phenylamino]-pyrimidin-4-yl}-1H-pyrazol-3-yl)-phenol;
[4-(4-methyl-piperazin-1-yl)-phenyl]-[4-(3-p-tolyl-1H-pyrazol-4-yl)-pyrimidin-2-yl]-amine;
[4-(2-dimethylamino-ethoxy)-phenyl]-[4-(3-p-tolyl-1H-pyrazol-4-yl)-pyrimidin-2-yl]-amine;
{4-[3-(4-chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[3-(1-methyl-piperidin-4-ylmethoxy)-phenyl]-amine;
{4-[3-(4-chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(1-methyl-piperidin-4-ylmethoxy)-phenyl]-amine;
{4-[3-(4-chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[3-(4-methyl-piperazin-1-yl)-phenyl]-amine;
4-{4-[3-(4-chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-ylamino}-benzoic acid;
(4-{4-[3-(4-chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-ylamino}-phenyl)-(4-methyl-piperazin-1-yl)-methanone;
{4-[3-(4-chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(4-methyl-piperazin-1-ylmethyl)-phenyl]-amine;
{4-[3-(2,4-dichloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-(3,4,5-trimethoxy-phenyl)-amine;
{4-[3-(2,4-dichloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(4-methyl-piperazin-1-yl)-phenyl]-amine;
N-{4-[3-(4-chloro-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-N',N'-dimethyl-benzene-1,3-diamine;
{4-[3-(4-ethoxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-[4-(4-methyl-piperazin-1-yl)-phenyl]-amine;
{4-[3-(4-ethoxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-(3,4,5-trimethoxy-phenyl)-amine;

{4-[3-(4-chloro-3-methoxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-(3,4,5-trimethoxy-phenyl)-amine;
3-{4-[3-(4-chloro-3-hydroxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-ylamino}-benzenesulfonamide;
3-{4-[3-(4-chloro-3-methoxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-ylamino}-benzenesulfonamide;
3-{4-[3-(4-chloro-3-methoxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-ylamino}-benzoic acid ethyl ester;
3-{4-[3-(4-chloro-3-hydroxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-ylamino}-benzoic acid ethyl ester;
1-(3-{4-[3-(4-chloro-3-methoxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-ylamino}-phenyl)-ethanone;
benzothiazol-6-yl-{4-[3-(4-chloro-3-methoxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-amine;
{4-[3-(4-chloro-3-methoxy-phenyl)-1H-pyrazol-4-yl]-pyrimidin-2-yl}-(1-oxo-benzothiazol-6-yl)-amine;
[4-(3-benzo[1,3]dioxol-5-yl-1H-pyrazol-4-yl)-pyrimidin-2-yl]-(3,4,5-trimethoxy-phenyl)-amine;
3-[4-(3-benzo[1,3]dioxol-5-yl-1H-pyrazol-4-yl)-pyrimidin-2-ylamino]-benzoic acid tert-butyl ester;
3-[4-(3-benzo[1,3]dioxol-5-yl-1H-pyrazol-4-yl)-pyrimidin-2-ylamino]-benzenesulfonamide;
[4-(3-benzo[1,3]dioxol-5-yl-1H-pyrazol-4-yl)-pyrimidin-2-yl]-(3-methylsulfanyl-phenyl)-amine;
and pharmaceutically acceptable salts thereof.

7. A compound of formula I, or a pharmaceutically acceptable salt thereof, according to any one of claims 1 to 6 for use in a method for the treatment of the human or animal body.

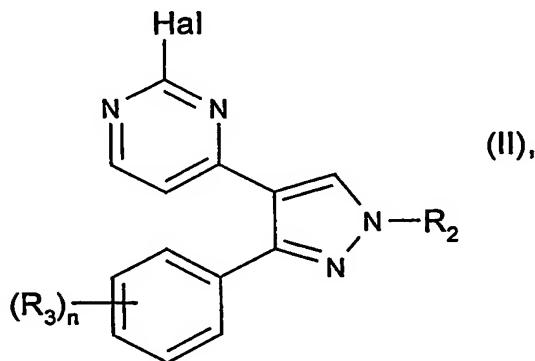
8. A pharmaceutical composition comprising a compound of formula I or a pharmaceutically acceptable salt thereof according to any one of claims 1 to 6, together with at least one pharmaceutically acceptable carrier.

9. Use of a compound of formula I according to any one of claims 1 to 6, or a pharmaceutically acceptable salt thereof, for the preparation of a pharmaceutical composition for the treatment of a disease.

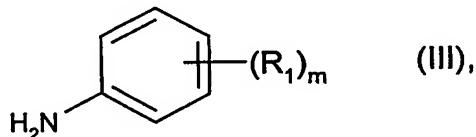
10. Use of a compound of formula I according to any one of claims 1 to 6, or a pharmaceutically acceptable salt thereof, for the preparation of a pharmaceutical composition for the treatment of a disease which responds to an inhibition of a protein tyrosine kinase.

11. A process for the preparation of a compound of formula I according to claim 1 or of a salt of such a compound, characterized in that

a) in order to prepare a compound of formula I, wherein R_2 is hydrogen or lower alkyl,
(i) a compound of formula II

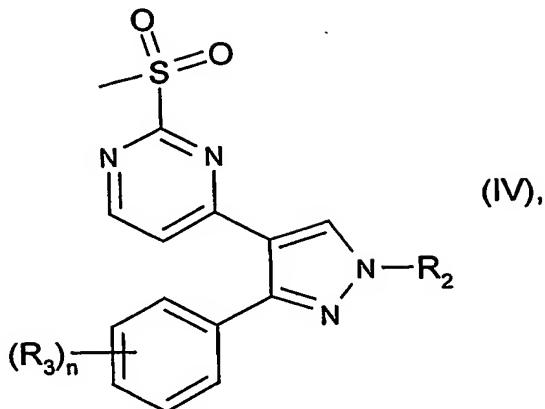


wherein Hal is halogen, R_2 is hydrogen or lower alkyl and n and R_3 have the meanings as defined for a compound of formula I, is reacted with a compound of formula III



wherein m and R_1 have the meanings as defined for a compound of formula I, or

(ii) a compound of formula IV



wherein R₂ is hydrogen or lower alkyl and n and R₃ have the meanings as defined for a compound of formula I, is reacted with a compound of formula III; or

b) in order to prepare a compound of formula I, wherein R₂ is unsubstituted or substituted lower alkyl or a heterocyclic radical, a compound of formula I, wherein R₂ is hydrogen, is reacted with a compound of the formula R₂-OH, wherein R₂ is unsubstituted or substituted lower alkyl or a heterocyclic radical wherein the unsubstituted or substituted lower alkyl or the heterocyclic radical is attached to the hydroxy group of R₂-OH via a carbon atom of the lower alkyl moiety or via a ring carbon atom of the heterocyclic radical, respectively;

whereby functional groups which are present in the starting compounds of process a) or b) and are not intended to take part in the reaction, are present in protected form if necessary, and protecting groups that are present are cleaved, whereby the said starting compounds may also exist in the form of salts provided that a salt-forming group is present and a reaction in salt form is possible;

and, if so desired, a compound of formula I thus obtained is converted into another compound of formula I, a free compound of formula I is converted into a salt, an obtained salt of a compound of formula I is converted into the free compound or another salt, and/or a mixture of isomeric compounds of formula I is separated into the individual isomers.

